

Access DB# 133  
135523

**SEARCH REQUEST FORM**

Scientific and Technical Information Center

Requester's Full Name: JANE ZARA Examiner #: 77512 Date: 10/19/04  
Art Unit: 1635 Phone Number: 302-0765 Serial Number: 09/802,376  
Mail Box and Bldg/Room Location: 2d28 Results Format Preferred (circle): PAPER DISK E-MAIL  
2018

If more than one search is submitted, please prioritize searches in order of need.

\*\*\*\*\*

Please provide a detailed statement of the search topic, and describe as specifically as possible the subject matter to be searched. Include the elected species or structures, keywords, synonyms, acronyms, and registry numbers, and combine with the concept or utility of the invention. Define any terms that may have a special meaning. Give examples or relevant citations, authors, etc, if known. Please attach a copy of the cover sheet, pertinent claims and abstract.

Title of Invention: La Formulations  
Inventors (please provide full names): Van Nest et al. CRFE

Earliest Priority Filing Date: 3/9/01

\*For Sequence Searches Only\* Please include all pertinent information (parent, child, divisional, or issued patent numbers) along with the appropriate serial number.

Please search Seq ID No: 1

- No size limit

- size limit to 100 NDS

Please include interference search.

1-22 ak

\*\*\*\*\*  
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	Type of Search	Vendors and cost where applicable
Searcher: <u>Beverly C 2528</u>	NA Sequence (#) _____	STN <u><input checked="" type="checkbox"/></u> _____
Searcher Phone #: _____	AA Sequence (#) _____	Dialog _____
Searcher Location: _____	Structure (#) _____	Questel/Orbit _____
Date Searcher Picked Up: _____	Bibliographic _____	Dr. Link _____
Date Completed: _____	Litigation _____	Lexis/Nexis _____
Searcher Prep & Review Time: _____	Fulltext _____	Sequence Systems _____
Clerical Prep Time: _____	Patent Family _____	WWW/Internet _____
Online Time: _____	Other _____	Other (specify) <u>CGN</u>

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Result No.	Score	Query Match	\$	ID			Description
				DB	Length	ID	
1	22	100.0	22	2	AAV32079	Nucleoid	
2	22	100.0	22	2	AAV80097	Immunomod	
3	22	100.0	22	2	AAV80103	Immunomod	
4	22	100.0	22	2	AAV80102	Immunomod	
5	22	100.0	22	2	AAx36624	ISS-ODN D	
6	22	100.0	22	3	AAx14467	Immunosti	
7	22	100.0	22	3	AAx38072	Immunosti	
8	22	100.0	22	3	AAx38071	Immunosti	
9	22	100.0	22	3	AAx38065	Immunosti	
10	22	100.0	22	3	AAx90458	CpG adjuv	
11	22	100.0	22	3	AAx96253	Sequence	
12	22	100.0	22	3	AAx25876	Immunomod	
13	22	100.0	22	3	AAx5876	Immunosti	
14	22	100.0	22	3	AAx64051	CpG motif	
15	22	100.0	22	4	AAx20403	Immunomod	
16	22	100.0	22	4	AAx43338	Immunomod	
17	22	100.0	22	4	AAx73439	Immunomod	
18	22	100.0	22	4	AAx75992	Immunomod	
19	22	100.0	22	4	AAx77040	Immunomod	
20	22	100.0	22	4	AAx29800	Cholera t	
21	22	100.0	22	4	AAx44109	5' termin	

CC conjugate synergistically boost the magnitude of the host immune response  
 CC against an antigen to a level greater than that the host immune response to  
 CC either the IMM, antigen or ISS-PN alone. These responses to ISS-PN/IMM  
 CC conjugates are particularly acute during the important early phase of the  
 CC host immune response to an antigen. The ISS-PN/IMM conjugates boost both  
 CC humoral (antibody) and cellular (Th1 type) immune responses of the host.  
 CC Thus, use of the method to boost the immune responsiveness of a host to  
 CC subsequent challenge by a sensitising antigen without immunisation avoids  
 CC the risk of Th2-mediated, immunisation-induced anaphylaxis by suppressing  
 CC IgE production in response to the antigen challenge. The conjugates can  
 CC also be used to combat pathogenic infection and to stimulate therapeutic  
 CC angiogenesis to treat conditions in which localised blood flow plays a  
 CC significant etiological role, e.g. retinopathies  
 XX  
 SQ Sequence 22 BP; 6 A; 3 C; 7 G; 6 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 2; Length 22;  
 Best Local Similarity 100.0%; Pred. No. 0.24;  
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TGACTGTGAACGTTTCGAGATGA 22  
 |||||  
 DB 1 TGACTGTGAACGTTTCGAGATGA 22

RESULT 2  
 AAV80097  
 ID AAV80097 standard; DNA; 22 BP.  
 AC AAV80097;  
 DT 12-MAR-1999 (first entry)  
 XX  
 DE Immunomodulatory oligo comprising an ISS sequence.  
 XX  
 KW Immunomodulatory; immunostimulatory; octanucleotide; immune regulation;  
 KW ISS: cancer; allergy; asthma; hepatitis B infection; papillomavirus;  
 KW human immunodeficiency virus; influenza; herpes; M. tuberculosis; ss;  
 KW B. pertussis; malaria; plasmodia; leishmania; trypanosoma; schistosoma.  
 XX  
 OS Synthetic.

PN WO9855495-A2.  
 PD 10-DEC-1998.  
 PF 05-JUN-1998; 98WO-US011578.  
 PR 06-JUN-1997; 97US-0048793P.  
 XX (DYNA-) DYNAVAX TECHNOLOGIES CORP.  
 XX  
 PI Schwartz D, Roman M, Dina D;  
 DR WPI; 1999-059898/05.  
 XX  
 PT Immunostimulatory oligonucleotides regulate the immune system - and  
 PT contain an immune-stimulating octanucleotide sequence; for treating  
 PT cancer, allergic and infectious diseases.  
 XX  
 PS Claim 5; Page 29; 63pp; English.

XX The invention relates to immunomodulatory oligonucleotides that comprise  
 CC at least 1 immunostimulatory octanucleotide sequence (ISS) where the ISS  
 CC sequences are selected from the group consisting of AACGTTCC, AACGTTCCG,  
 CC GACGTTCC, and GACGTTCCG. The immunomodulatory sequences are used to treat  
 CC patients needing immune regulation, such as those suffering from cancer,  
 CC an allergic disease and asthma. They are also used to prevent infectious  
 CC diseases such as influenza, herpes, hepatitis B, human immunodeficiency  
 CC and papillomavirus, Hemophilus influenza, Mycobacterium tuberculosis and  
 CC Bordetella pertussis, malarial plasmodia, Leishmania, Trypanosoma and  
 CC Schistosoma. The immunomodulatory sequences are used to screen for human  
 CC immunostimulatory activity by incubating macrophage cells and the

CC oligonucleotide; and determining the relative amount of Th1-biased  
 CC cytokines in the supernatant. Sequences AAV80096 to AAV80103 represent  
 CC specific claimed examples of such immunomodulatory oligonucleotides  
 XX  
 SQ Sequence 22 BP; 6 A; 3 C; 7 G; 6 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 2; Length 22;  
 Best Local Similarity 100.0%; Pred. No. 0.24;  
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TGACTGTGAACGTTTCGAGATGA 22  
 |||||  
 DB 1 TGACTGTGAACGTTTCGAGATGA 22

RESULT 3  
 AAV80103  
 ID AAV80103 standard; DNA; 22 BP.  
 AC AAV80103;  
 DT 12-MAR-1999 (first entry)  
 XX  
 DE Immunomodulatory oligo comprising an ISS sequence.  
 XX  
 KW Immunomodulatory; immunostimulatory; octanucleotide; immune regulation;  
 KW ISS: cancer; allergy; asthma; hepatitis B infection; papillomavirus;  
 KW human immunodeficiency virus; influenza; herpes; M. tuberculosis; ss;  
 KW B. pertussis; malaria; plasmodia; leishmania; trypanosoma; schistosoma.  
 XX  
 OS Synthetic.

FH Key Location/Qualifiers  
 FT modified\_base 11 /tag= a  
 FT /note= "5-bromocytosine"

PN WO9855495-A2.  
 PD 10-DEC-1998.  
 PF 05-JUN-1998; 98WO-US011578.  
 PR 06-JUN-1997; 97US-0048793P.  
 XX (DYNA-) DYNAVAX TECHNOLOGIES CORP.  
 XX  
 PI Schwartz D, Roman M, Dina D;  
 DR WPI; 1999-059898/05.  
 XX  
 PT Immunostimulatory oligonucleotides regulate the immune system - and  
 PT contain an immune-stimulating octanucleotide sequence; for treating  
 PT cancer, allergic and infectious diseases.  
 XX  
 PS Claim 24; Page 30; 63pp; English.

XX The invention relates to immunomodulatory oligonucleotides that comprise  
 CC at least 1 immunostimulatory octanucleotide sequence (ISS) where the ISS  
 CC sequences are selected from the group consisting of AACGTTCC, AACGTTCCG,  
 CC GACGTTCC, and GACGTTCCG. The immunomodulatory sequences are used to treat  
 CC patients needing immune regulation, such as those suffering from cancer,  
 CC an allergic disease and asthma. They are also used to prevent infectious  
 CC diseases such as influenza, herpes, hepatitis B, human immunodeficiency  
 CC and papillomavirus, Hemophilus influenza, Mycobacterium tuberculosis and  
 CC Bordetella pertussis, malarial plasmodia, Leishmania, Trypanosoma and  
 CC Schistosoma. The immunomodulatory sequences are used to screen for human  
 CC immunostimulatory activity by incubating macrophage cells and the  
 CC oligonucleotide; and determining the relative amount of Th1-biased  
 CC cytokines in the supernatant. Sequences AAV80096 to AAV80103 represent  
 CC specific claimed examples of such immunomodulatory oligonucleotides  
 XX  
 SQ Sequence 22 BP; 6 A; 3 C; 7 G; 6 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 2; Length 22;  
 Best Local Similarity 100.0%; Pred. No. 0.24;  
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGAAGTGTGAACGTTTCGAGATGA 22  
 DB 1 TGAAGTGTGAACGTTTCGAGATGA 22

## RESULT 4

AAV80102

ID AAV80102 standard; DNA; 22 BP.

XX

AC AAV80102;

XX

DT 12-MAR-1999 (first entry)

XX

DE Immunomodulatory oligo comprising an ISS sequence.

XX

KW Immunomodulatory; immunostimulatory; octanucleotide; immune regulation;

KW ISS: cancer; allergy; asthma; hepatitis B infection; papillomavirus;

KW human immunodeficiency virus; influenza; herpes; M. tuberculosis; ss;

KW B. pertussis; malaria; plasmodia; Leishmania; Trypanosoma; Schistosoma.

XX

OS Synthetic.

XX

FH Key Location/Qualifiers

XX modified\_base 11

FT /\*tag= a

FT /note= "5-bromocytosine"

FT

XX WO9855495-A2.

XX

XX 10-DEC-1998.

XX

XX 05-JUN-1998; 98WO-US011578.

XX

XX 06-JUN-1997; 97US-0048793P.

XX

XX (DYNA-) DYNAVAX TECHNOLOGIES CORP.

XX

XX Schwartz D, Roman M, Dina D;

XX

XX WPI; 1999-059898/05.

XX

XX Immunostimulatory oligonucleotides regulate the immune system - and

XX contain an immune-stimulating octanucleotide sequence; for treating

XX cancer, allergic and infectious diseases.

XX

XX Claim 23; Page 30; 63pp; English.

XX

XX The invention relates to immunomodulatory oligonucleotides that comprise

XX at least 1 immunostimulatory octanucleotide sequence (ISS) where the ISS

XX sequences are selected from the group consisting of AAGTTCC, AAGTTCCG,

XX CCAGTTCC, and GAGTTCC. The immunomodulatory sequences are used to treat

XX patients needing immune regulation, such as those suffering from cancer,

XX an allergic disease and asthma. They are also used to prevent infectious

XX diseases such as influenza, herpes, hepatitis B, human immunodeficiency

XX and papillomavirus, Hemophilus influenza, Mycobacterium tuberculosis and

XX Bordetella pertussis, malarial plasmodia, Leishmania, Trypanosoma and

XX Schistosoma. The immunomodulatory sequences are used to screen for human

XX immunostimulatory activity by incubating macrophage cells and the

XX oligonucleotide; and determining the relative amount of Th1-biased

XX cytokines in the supernatant. Sequences AAV80096 to AAV80103 represent

XX specific claimed examples of such immunomodulatory oligonucleotides

XX

XX Sequence 22 BP; 6 A; 3 C; 7 G; 6 T; 0 U; 0 Other;

XX

Query Match 100.0%; Score 22; DB 2; Length 22;

Best Local Similarity 100.0%; Pred. No. 0.24;

Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGAAGTGTGAACGTTTCGAGATGA 22  
 DB 1 TGAAGTGTGAACGTTTCGAGATGA 22

## RESULT 5

AAV36624

ID AAV36624 standard; DNA; 22 BP.

XX

AC AAV36624;

XX

DT 09-JUL-1999 (first entry)

XX

DE ISS-ODN DY1018 nucleotide sequence.

XX

KW Antigen-stimulated inflammation; immunostimulatory oligonucleotide;

KW granulocyte-mediated tissue inflammation; Th2 type immune response;

KW immune responsiveness modulation; idiopathic hypersensitivity syndrome;

KW cutaneous basophil hypersensitivity; ISS-ODN; asthma; nasal polypsis;

KW allergic rhinitis; atopic dermatitis; allergic conjunctivitis;

KW eosinophilic fasciitis; therapy; ss.

XX

OS Synthetic.

XX

PN WO9911275-A2.

XX

XX 11-MAR-1999.

XX

XX 04-SEP-1998; 98WO-US018382.

XX

XX 05-SEP-1997; 97US-00927120.

XX

XX (REGC ) UNIV CALIFORNIA.

XX

XX Ray E;

XX

XX WPI; 1999-312404/26.

XX

XX Reducing antigen-stimulated granulocyte-mediated inflammation.

XX

XX Example 2; Page 30; 69pp; English.

XX

XX This is the ISS-ODN DY1018 nucleotide sequence. The invention relates to

XX a method for preventing or reducing antigen-stimulated, granulocyte-

XX mediated tissue inflammation in a mammal, by administering an

XX immunostimulatory oligonucleotide (ISS-ODN), where: (a) reduction in, or

XX the absence of, a Th2 type immune response is measured; or (b) there is a

XX reduction or absence of other clinical signs of inflammation in the host

XX after antigen challenge. The method is used to reduce or suppress

XX granulocyte-mediated inflammation in a host tissue, and to modulate the

XX host's immune responsiveness to an antigen, particularly where the

XX subject suffers from asthma, nasal polypsis, allergic rhinitis, atopic

XX dermatitis, allergic conjunctivitis, eosinophilic fasciitis, idiopathic

XX hypersensitivity syndrome, or cutaneous basophil hypersensitivity.

XX Unlike prior art treatment by antigen immunisation, the method is an

XX antigen-independent method, and avoids host production of both

XX interleukin-4 (IL-4), which carries risk of anaphylaxis, and IL-5 which

XX actually encourages granulocyte adhesion to endothelia

XX

XX Sequence 22 BP; 6 A; 3 C; 7 G; 6 T; 0 U; 0 Other;

XX

Query Match 100.0%; Score 22; DB 2; Length 22;

Best Local Similarity 100.0%; Pred. No. 0.24;

Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGAAGTGTGAACGTTTCGAGATGA 22  
 DB 1 TGAAGTGTGAACGTTTCGAGATGA 22

## RESULT 6

AAV14467

ID AAV14467 standard; DNA; 22 BP.

XX AAA14467;  
 XX  
 XX  
 XX  
 XX 21-AUG-2000 (first entry)  
 XX  
 XX Immunostimulatory oligonucleotide (ISS-ODN) DY1018.  
 DE  
 XX Immunostimulatory oligonucleotide; adjuvant; mucosal immunity;  
 KW  
 XX secretory immunoglobulin A production; sigA; Th1 phenotype; ds.  
 KW  
 XX  
 XX Synthetic.  
 OS  
 XX  
 XX Key modified\_base 11 Location/Qualifiers  
 PN /tag= a  
 XX /mod\_base= OTHER  
 XX 13-APR-2000.  
 PD /note= "5-Bromocytosine"  
 XX  
 XX 15-SEP-1999; 99WO-US021203.  
 PF  
 XX 05-OCT-1998; 98US-00167039.  
 PR  
 XX (REGC ) UNIV CALIFORNIA.  
 PA  
 XX Raz E, Horner AA, Carson DA;  
 PI  
 XX WPI; 2000-303647/26.  
 DR  
 XX Immunostimulatory oligonucleotide adjuvant induces mucosal immunity to an  
 PT antigen in a mammalian host through production of secretory  
 PT immunoglobulin A.  
 PT  
 XX Claim 8; Page 21; 64pp; English.  
 PS  
 XX The invention relates to a method of inducing mucosal immunity to an  
 CC antigen in a mammalian host, including the production of secretory  
 CC immunoglobulin A (sIgA). Immune protection in the mucosa (the principal  
 CC site of entry of most foreign antigens) is mediated by mucosa-associated  
 CC lymphoid tissue, epithelial and distinct B-cell, T-cell and accessory  
 CC cell sub-populations. The primary immune response which characterises the  
 CC induction of mucosal immunity to an antigen is sIgA production by  
 CC activated B-cells. The method comprises introducing an immunostimulatory  
 CC oligonucleotide (ISS-ODN) and the antigen into host mucosa, where the ISS  
 CC -ODN includes a core nucleotide sequence. The core nucleotide sequence is  
 CC 5'-Purine-Purine-C-G-Pyrimidine-Pyrimidine-3', specific examples of which  
 CC are AACGTT, AGCGTC and GACGTT (SEQ ID NOS 1-3). A specific example of an  
 CC ISS-ODN is DY1018 (AAA14467). The ISS-ODN is used as an adjuvant with an  
 CC antigen for stimulating mucosal immunity. The level of sIgA production  
 CC induced in the host is at least 3 times the magnitude of sIgA production  
 CC achievable in response to introduction of antigen alone into the mucosal  
 CC tissue and is equivalent or greater than the magnitude of sIgA production  
 CC achievable in response to introduction of the antigen and cholera toxin  
 CC adjuvant into the mucosal tissue. The host immune response is stimulated  
 CC to antigen specific IgA production, biased towards the Th1 phenotype  
 CC while antigen-induced IgE production is avoided. The adjuvant has little  
 CC or no known toxicity in mammals and its efficacy is comparable to that of  
 CC cholera toxin which is used as a mucosal adjuvant. The present sequence  
 CC represents the immunostimulatory oligonucleotide DY1018  
 XX  
 XX Sequence 22 BP; 6 A; 3 C; 7 G; 6 T; 0 U; 0 Other;  
 SQ  
 Query Match 100.0%; Score 22; DB 3; Length 22;  
 Best Local Similarity 100.0%; Pred. NO. 0.24;  
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 TGACTGTGAACGTTTCGAGATGA 22  
 DB 1 TGACTGTGAACGTTTCGAGATGA 22  
 RESULT 7  
 AAA38072  
 ID AAA38072 standard; DNA; 22 BP.  
 XX  
 AC AAA38072;

XX 24-AUG-2000 (first entry)  
 XX  
 XX Immunostimulatory sequence (ISS) #7.  
 XX  
 XX Immunostimulatory sequence; ISS; immunomodulator; glycoprotein 120;  
 KW gp120; human immunodeficiency virus; HIV; immune response; infection;  
 KW development; ss.  
 KW  
 XX Synthetic.  
 OS  
 XX  
 XX Key modified\_base 11 Location/Qualifiers  
 PN /tag= a  
 XX /mod\_base= OTHER  
 XX 13-APR-2000.  
 PD /note= "5-Bromocytosine"  
 XX  
 XX 15-SEP-1999; 99WO-US023677.  
 PF  
 XX 09-OCT-1998; 98US-0103733P.  
 PR  
 XX 07-OCT-1999; 99US-00415186.  
 XX  
 XX (DYNA-) DYNAVAX TECHNOLOGIES CORP.  
 PA  
 XX Tighe H, Raz E, Schwartz D, Takabayashi K;  
 PI  
 XX WPI; 2000-317846/27.  
 DR  
 XX Anti-HIV composition comprises immunostimulatory polynucleotides and HIV  
 PT glycoprotein gp120 useful for modulating, stimulating an immune response  
 PT against HIV in an HIV infected individual.  
 PT  
 XX Disclosure; Page 17; 65pp; English.  
 PS  
 XX The present invention relates to an immunostimulatory composition  
 CC comprising a human immunodeficiency virus (HIV) antigen, and an  
 CC immunomodulatory polynucleotide comprising an immunostimulatory sequence  
 CC (ISS). This sequence represents an ISS that can be used in the  
 CC composition. An immunostimulatory composition which comprises a gp120  
 CC conjugated to an immunomodulatory polynucleotide, or is proximately  
 CC associated to it and not conjugated, is used for modulating or  
 CC stimulating a specific immune response against gp120 in an individual by  
 CC producing anti-gp120 antibodies or gp120 specific cytotoxic T cells. It  
 CC is also used for suppressing or delaying development of HIV infection in  
 CC an individual infected with HIV or an individual at risk of infection in  
 CC with HIV, respectively. It is also used for treating an individual  
 CC infected with HIV in need of immune modulation  
 XX  
 XX Sequence 22 BP; 6 A; 3 C; 7 G; 6 T; 0 U; 0 Other;  
 SQ  
 Query Match 100.0%; Score 22; DB 3; Length 22;  
 Best Local Similarity 100.0%; Pred. NO. 0.24;  
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 TGACTGTGAACGTTTCGAGATGA 22  
 DB 1 TGACTGTGAACGTTTCGAGATGA 22  
 RESULT 8  
 AAA38071  
 ID AAA38071 standard; DNA; 22 BP.  
 XX  
 AC AAA38071;  
 XX

```

DT 24-AUG-2000 (first entry)
DE Immunostimulatory sequence (ISS) #7.
KW Immunostimulatory sequence; ISS; immunomodulator; glycoprotein 120;
KW gp120; human immunodeficiency virus; HIV; immune response; infection;
KW development; ss.
XX
OS Synthetic.
XX
FH Key Location/Qualifiers
FT modified_base 11
FT /*tag= a
FT /mod_base= OTHER
FT /note= "5-Bromocytosine"
XX
PN WO200021556-A1.
XX
XX 20-APR-2000.
XX
PF 08-OCT-1999; 99WO-US023677.
XX
PR 09-OCT-1998; 98US-0103733P.
XX
PR 07-OCT-1999; 99US-00415186.
XX
XX (DYNA-) DYNAVAX TECHNOLOGIES CORP.
XX
XX Tighe H, Raz E, Schwartz D, Takabayashi K;
XX WPI; 2000-317846/27.
XX
XX Anti-HIV composition comprises immunostimulatory polynucleotides and HIV
XX glycoprotein gp120 useful for modulating, stimulating an immune response
XX against HIV in an HIV infected individual.
XX
XX Disclosure; Page 17; 65pp; English.
XX
XX The present invention relates to an immunostimulatory composition
XX comprising a human immunodeficiency virus (HIV) antigen, and an
XX immunomodulatory polynucleotide comprising an immunostimulatory sequence
XX (ISS). This sequence represents an ISS that can be used in the
XX composition. An immunostimulatory composition which comprises a gp120
XX conjugated to it and not conjugated, is used for modulating or
XX stimulating a specific immune response against gp120 in an individual by
XX producing anti-gp120 antibodies or gp120 specific cytotoxic T cells. It
XX is also used for suppressing or delaying development of HIV infection in
XX an individual infected with HIV or an individual at risk of infection
XX with HIV, respectively. It is also used for treating an individual
XX infected with HIV in need of immune modulation
XX
SQ Sequence 22 BP; 6 A; 3 C; 7 G; 6 T; 0 U; 0 Other;
Query Match 100.0%; Score 22; DB 3; Length 22;
Best Local Similarity 100.0%; Pred. No. 0.24;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TGACTGTGAACGTCGAGATGA 22
DB 1 TGACTGTGAACGTCGAGATGA 22
RESULT 10
AAA90458
ID AAA90458 standard; DNA; 22 BP.
XX
AC AAA90458;
XX
XX 10-JAN-2001 (first entry)
XX
DE Cpg adjuvant oligonucleotide, SEQ ID NO:19.
XX
XX Cpg oligonucleotide; CpG motif; adjuvant; microdroplet emulsion;
KW microemulsion; adsorbent microparticle; vaccine; Th1 immune response;
KW viral infection; bacterial infection; parasitic infection; HCV; HBV;
KW hepatitis C virus; hepatitis B virus; herpes simplex virus; HSV; HIV;
KW human immunodeficiency virus; cytomegalovirus; CMV; influenza virus;
KW rabies virus; cholera; diphtheria; tetanus; pertussis;
KW Helicobacter pylori; Haemophilus influenzae; malaria; ss.
XX
OS Synthetic.
XX
PN WO200050006-A2.
XX

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PD 31-AUG-2000.
XX
XX 09-FEB-2000; 2000WO-US003331.
XX
XX 26-FEB-1999; 99US-0121858P.
PR 29-JUL-1999; 99US-0146391P.
PR 28-OCT-1999; 99US-0161997P.
XX
XX (CHIR ) CHIRON CORP.
PA
XX O'hagan D, Ott GS, Donnelly J, Kazzaz J, Uguzzoli M, Singh M;
PI Barackman J;
PI
XX WPI; 2000-587123/55.
DR
XX Microemulsion having an adsorbent surface comprising a microdroplet
PT emulsion consisting of a metabolizable oil and an emulsifying agent which
PT is a detergent, useful as a vaccine to treat bacterial, viral, and
PT parasitic infection.
XX
XX Claim 17; Page 40; 95pp; English.
PS
XX The invention relates to a microdroplet emulsion (microemulsion) with an
CC adsorbent surface, and which comprises a metabolizable oil and an
CC emulsifying agent (a detergent). It also relates to a composition
CC comprising the microemulsion and a microparticle with an adsorbent
CC surface, where the microparticle comprises a polymer selected from a
CC poly(alpha-hydroxy acid), a polyhydroxy butyric acid, a polycaprolactone,
CC a polyorthoester, a polyanhydride, and a polycyanoacrylate, and a second
CC detergent. The surface of the microparticles efficiently adsorb
CC biologically active macromolecules such as DNA, polypeptides, antigens,
CC hormones, pharmaceuticals, enzymes, mediators of transcription or
CC translation, metabolic intermediates and adjuvants. Additionally, a
CC second biologically active molecule may be encapsulated within the
CC microparticle. The microemulsion can be used in methods of immunising a
CC host animal, particularly a human, against a viral, bacterial or
CC parasitic infection, and in methods of increasing a Th1 immune response.
CC The microemulsions (having the appropriate antigens adsorbed) may be
CC particularly used as vaccines for hepatitis C virus (HCV), hepatitis B
CC virus (HBV), herpes simplex virus (HSV), human immunodeficiency virus
CC (HIV), cytomegalovirus (CMV), influenza virus, and rabies virus; the
CC bacteria which cause cholera, diphtheria, tetanus and pertussis;
CC Helicobacter pylori and Haemophilus influenzae; and malaria-causing
CC parasites. Sequences AAA90447-A90467 represent Th1 lymphocyte stimulating
CC oligonucleotides containing at least one CpG motif which are claimed for
CC use as adjuvants in the compositions of the invention
XX
XX Sequence 22 BP; 6 A; 3 C; 7 G; 6 T; 0 U; 0 Other;
SQ
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Best Local Similarity 100.0%; Pred. No. 0.24; Mismatches 0; Gaps 0;
Matches 22; Conservative 0; Indels 0;
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Db 1 TGAAGTGTGAACGTTTCGAGATGA 22
RESULT 12
AAA96253
ID AAA96253 standard; DNA; 22 BP.
XX
XX AAA96253;
AC
XX
XX 08-FEB-2001 (first entry)
DT
XX
XX Sequence of a stabilised oligonucleotide with antitumour activity.
DE
XX Antitumour; immunostimulatory oligonucleotide; tumour; anaplasia;
KW glioblastoma; medullablastoma; neuroblastoma; carcinoma; ss.
KW
XX
XX Synthetic.
OS
XX
Query Match 100.0%; Score 22; DB 3; Length 22;
Best Local Similarity 100.0%; Pred. No. 0.24; Mismatches 0; Gaps 0;
Matches 22; Conservative 0; Indels 0;
QY 1 TGAAGTGTGAACGTTTCGAGATGA 22
Db 1 TGAAGTGTGAACGTTTCGAGATGA 22
RESULT 11
AAA96253
ID AAA96253 standard; DNA; 22 BP.
XX
XX AAA96253;
AC
XX
XX 08-FEB-2001 (first entry)
DT
XX
XX Sequence of a stabilised oligonucleotide with antitumour activity.
DE
XX Antitumour; immunostimulatory oligonucleotide; tumour; anaplasia;
KW glioblastoma; medullablastoma; neuroblastoma; carcinoma; ss.
KW
XX
XX Synthetic.
OS
XX

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PN WO200056342-A2.
XX
XX 28-SEP-2000.
XX
XX 17-MAR-2000; 2000WO-FR000676.
XX
XX 19-MAR-1999; 99FR-00003433.
XX
XX (ASSI-) ASSISTANCE PUBLIQUE HOPITAUX PARIS.
PA (INRM ) INST NAT SANTE & RECH MEDICALE.
XX
XX Carpentier A;
PI
XX WPI; 2000-602192/57.
DR
XX Use of stabilized oligonucleotides as antitumor agents, particularly
PT against nervous system tumors, have optimal activity and are not toxic.
PT
XX Example 2; Page 16; 57pp; French.
PS
XX The present sequence represents a stabilised oligonucleotide which has
CC antitumour activity. The oligonucleotide comprises an octamer motif of
CC the type 5'-purine-purine-CG-pyrimidine-pyrimidine-X-X-3', where the pair
CC X-X is AT, AA, CT or TT. The oligonucleotides are immunostimulatory, and
CC are not toxic. They may be adapted for use in animals or humans. The
CC stabilised oligonucleotides are used for treating tumours, of any type
CC and any degree of anaplasia, particularly human tumours in the peripheral
CC or central nervous systems, specifically glioblastomas, medullablastomas,
CC neuroblastomas, melanomas or carcinomas
XX
XX Sequence 22 BP; 6 A; 3 C; 7 G; 6 T; 0 U; 0 Other;
SQ
Query Match 100.0%; Score 22; DB 3; Length 22;
Best Local Similarity 100.0%; Pred. No. 0.24; Mismatches 0; Gaps 0;
Matches 22; Conservative 0; Indels 0;
QY 1 TGAAGTGTGAACGTTTCGAGATGA 22
Db 1 TGAAGTGTGAACGTTTCGAGATGA 22
RESULT 12
AAZ55876
ID AAZ55876 standard; DNA; 22 BP.
XX
XX AAZ55876;
AC
XX
XX 10-APR-2000 (first entry)
DT
XX
XX Immunomodulatory oligonucleotide SEQ ID NO: 1.
DE
XX Immunomodulation; immunostimulatory sequence; adjuvant;
KW Th1 immune response; cytotoxic T-cell; cytokine; cancer; allergy; asthma;
KW immunosuppression; ss.
XX
XX Mus musculus.
OS
XX Synthetic.
XX
XX Key Location/Qualifiers
FH modified_base 1..22
FT /tag= a
FT /note= "Phosphorothioate linkages"
FT misc_feature 9..16
FT /tag= b
FT /note= "Immunostimulatory sequence (ISS)"
XX
XX WO962923-A2.
PN
XX
XX 09-DEC-1999.
PD
XX
XX 04-JUN-1999; 99WO-US012538.
PF
XX
XX 05-JUN-1998; 98US-0088310P.
PR

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PR 01-JUN-1999; 99US-00324191.
XX (DYNA-) DYNAVAX TECHNOLOGIES CORP.
PA Schwartz D;
XX WPI; 2000-105687/09.
XX Novel immunomodulatory oligonucleotide used to induce a Th1-type immune
PT response, e.g. to tumor antigens.
XX Example 1; Page 35; 54pp; English.
XX Sequences AAZ55876-255877 and AAZ55880-255886 represent immunomodulatory
CC oligonucleotides comprising an immunostimulatory sequence (ISS, e.g.,
CC AACGTC, AACGTC, AGCGTC, AGCGTT, GACGTC, GACGTT, GCGGTT, AACGTTTC
CC and GACGTTTC). The invention relates to oligonucleotides comprising one
CC or more ISSs, where the ISS comprises at least one modified cytosine with
CC an electron-withdrawing moiety at position C-5 or C-6 of the base.
CC Sequences AAZ55877 and AAZ55880-255886 contain ISSs comprising at least
CC one bromocytosine, whereas sequence AAZ55876 contains an unmodified ISS.
CC The immunomodulatory oligonucleotides have an adjuvant-like effect; when
CC formulated with an antigen, the oligonucleotides stimulate production of
CC Th1-type cytokines, and induce a Th1-type immune response (activation of
CC cytotoxic T cells), while simultaneously downregulating the Th2-type
CC response. The Th1 response is particularly effective for control of
CC viruses and intracellular parasites. The immunomodulatory
CC oligonucleotides are used, particularly when formulated with an antigen
CC or a facilitator, for modulating immune responses. Such compositions may
CC be used in tumour therapy, in treatment of allergy (including asthma),
CC for inducing a vigorous cellular response (against a virus, bacterium,
CC fungus or protozoan), and also in contraceptive vaccines based on sperm
CC antigens
XX Sequence 22 BP; 6 A; 3 C; 7 G; 6 T; 0 U; 0 Other;
SQ Query Match 100.0%; Score 22; DB 3; Length 22;
Best Local Similarity 100.0%; Pred. No. 0.24;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TGACTGTGAACGTTTCGAGATGA 22
DB 1 TGACTGTGAACGTTTCGAGATGA 22

RESULT 13
AAC64051
ID AAC64051 standard; DNA; 22 BP.
XX AAC64051;
XX 15-FEB-2001 (first entry)
XX Immunostimulatory CpG phosphorothioate oligodeoxynucleotide.
DE CpG oligodeoxynucleotide; phosphorothioate; immunostimulatory; ISS ODN;
XX enhanced antigen presentation; antigen-presenting cell; APC;
KW T-cell activation; tumour cell; tumour antigen; cancer immunotherapy;
KW vaccine; ss.
XX Synthetic.
OS WO200062787-A1.
XX PN WO200062787-A1.
XX PD 26-OCT-2000.
XX PF 11-APR-2000; 2000WO-US009664.
XX PR 15-APR-1999; 99US-00292278.
XX (REGC ) UNIV CALIFORNIA.
PA Raz E, Martin-Orozco E;
PI

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XX WPI; 2000-679548/66.
XX Enhancing antigen-presentation capabilities of T-cells for cancer
PT immunotherapy, by contacting cells with an immunostimulatory
PT oligonucleotide.
XX Example 1; Page 18; 42pp; English.
XX The invention relates to a method of inducing activation of T-cells to
CC respond to an antigen, comprising contacting antigen-presenting cells
CC (APC) with an immunostimulatory oligodeoxynucleotide (ISS-ODN). The APCs
CC thus treated have enhanced antigen presenting capabilities compared to
CC antigen-activated APCs. APCs with enhanced antigen-presentation
CC capabilities then present the antigen to T-cells. The method is useful
CC for cancer immunotherapy. The ISS-ODN is used to enhance the tumour
CC antigen presenting capacity of tumour cells, thereby inducing T-cell
CC activation, and is therefore useful for treating tumours. Additionally,
CC tumour cells treated with an ISS-ODN ex vivo are useful as vaccines. ISS-
CC ODN treated APCs are induced to take up antigen through upregulation of
CC Fc-receptor expression, to present antigen through upregulation of major
CC histocompatibility complex (MHC) Class I and II expression and CD1d
CC expression, to produce co-stimulatory factors (B7 and CD40), to provide
CC cell-to-cell adhesion through upregulation of intercellular adhesion
CC molecule (ICAM) expression, and to increase Th1 stimulatory cytokine
CC production, all at levels greater than that achieved through contact of
CC APC with antigen alone. The present sequence represents a
CC phosphorothioate CpG ISS-ODN used in the exemplifications of the
CC invention
XX Sequence 22 BP; 6 A; 3 C; 7 G; 6 T; 0 U; 0 Other;
SQ Query Match 100.0%; Score 22; DB 3; Length 22;
Best Local Similarity 100.0%; Pred. No. 0.24;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TGACTGTGAACGTTTCGAGATGA 22
DB 1 TGACTGTGAACGTTTCGAGATGA 22

RESULT 14
AAH20403
ID AAH20403 standard; DNA; 22 BP.
XX AAH20403;
XX 03-AUG-2001 (first entry)
XX CpG motif containing oligonucleotide SEQ ID #21.
XX Immune system stimulator; CpG motif; CpG receptor; CpG-R; antibacterial;
KW immune response; vaccine adjuvant; tumour immunotherapy; allergy;
KW anti-inflammatory; cystic fibrosis; sepsis; heart disease; chlamydia;
KW inflammatory bowel disease; arthritis; multiple sclerosis; ss.
XX Unidentified.
XX Key Location/Qualifiers
FT modified_base 1...22 /*tag= a
FT /mod_base= OTHER
FT /note= "Phosphorothioate internucleoside linkages"
XX WO200132877-A2.
XX PD 10-MAY-2001.
XX PF 01-NOV-2000; 2000WO-US041735.
XX PR 02-NOV-1999; 99US-0163157P.
XX 24-NOV-1999; 99US-0167389P.
XX

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PA (CHIR ) CHIRON CORP.  
 XX Mackichan ML;  
 XX WPI; 2001-343486/36.  
 DR  
 XX Novel CpG receptor and nucleic acid molecule encoding the receptor, for  
 PT modulating immune response and for identifying compounds of therapeutic  
 PT use which bind and/or modulate the activity of the receptor.  
 XX  
 PS Example 1; Page 14; 41pp; English.  
 XX  
 CC Unmethylated CG dinucleotide sequences are commonly found in bacterial  
 CC DNA, and have been found to stimulate the innate immune system. Natural  
 CC killer and T cells are activated by exposure to oligonucleotides  
 CC containing CpG motifs. Oligonucleotides containing CpG motifs can be used  
 CC as adjuvants in vaccines. The present invention relates to a CpG  
 CC receptor. The CpG receptor contains a Toll homology domain (THD). The  
 CC Toll receptor family are associated with responses to pathogens. CpG  
 CC oligonucleotides may act as stimulators of various immune responses. The  
 CC CpG receptor or cells expressing the receptor are useful for identifying  
 CC a compound which binds to or modulates an activity of the CpG receptor.  
 CC The compounds are useful in e.g. vaccine adjuvants promoting cell-  
 CC mediated immune responses, antibacterials, (e.g. protection from *Listeria*  
 CC infection), tumour immunotherapy, allergy treatment, (e.g. suppressing  
 CC IgE in human PBMC, shifting from Th2 to Th1) and as anti-inflammatory  
 CC agents (e.g. for use in cystic fibrosis, sepsis, heart disease,  
 CC chlamydia, inflammatory bowel disease, arthritis and multiple sclerosis).  
 CC The present sequence represents a CpG motif containing oligonucleotide  
 CC used in examples demonstrating that CpG oligonucleotides can activate the  
 CC MAPK pathways and NF-kappaB  
 XX  
 SQ Sequence 22 BP; 6 A; 3 C; 7 G; 6 T; 0 U; 0 Other;  
 Query Match 100.0%; Score 22; DB 4; Length 22;  
 Best Local Similarity 100.0%; Pred. No. 0.24;  
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 OY 1 TGACTGTGAACGTTTCGAGATGA 22  
 DB 1 TGACTGTGAACGTTTCGAGATGA 22  
 RESULT 15  
 AAH43338  
 ID AAH43338 standard; DNA; 22 BP.  
 XX  
 AC AAH43338;  
 XX  
 DT 13-DEC-2001 (first entry)  
 DE Immunomodulatory polynucleotide 1018.  
 XX  
 KW Immunomodulation; inflammation; gastrointestinal tract;  
 KW ulcerative colitis; Crohn's disease; inflammatory bowel disease;  
 KW diarrhoea; rectal bleeding; weight loss; colon; weight; lesion; ss.  
 XX  
 OS Synthetic.  
 XX  
 PN WO200162207-A2.  
 XX  
 PD 30-AUG-2001.  
 XX  
 PF 22-FEB-2001; 2001WO-US006034.  
 XX  
 PR 23-FEB-2000; 2000US-0184256P.  
 XX  
 PA (REGC ) UNIV CALIFORNIA.  
 XX  
 PI Raz E, Rachmillewitz D;  
 XX  
 DR WPI; 2001-565393/63.  
 XX

PT Ameliorating gastrointestinal inflammation e.g. inflammatory bowel  
 PT disease involves administering an immunomodulatory nucleic acid.  
 XX  
 PS Claim 7; Page 28; 58pp; English.  
 XX  
 CC The sequences given in AAH43338-48 represent immunomodulatory  
 CC polynucleotides which may be used to ameliorate inflammation of the  
 CC gastrointestinal tract by administering a nucleic acid comprising one of  
 CC these sequences. These polynucleotides all comprise an immunomodulatory  
 CC nucleotide sequence of 5'-CpG-3' (1). The nucleotides may be used for  
 CC ameliorating or reducing gastrointestinal inflammation e.g. chronic or  
 CC acute gastrointestinal inflammation, ulcerative colitis, Crohn's disease  
 CC caused by inflammatory bowel disease; diarrhoea, rectal bleeding, weight  
 CC loss; to reduce colon weight and colon lesions; to reduce a colonic  
 CC inflammation. The immunomodulatory polynucleotides treat inflammatory  
 CC bowel disease satisfactorily and effectively and have little or no  
 CC toxicity even at a high dosage of 50000 micro-g. They also reduce the  
 CC risk of colonic cancer by treating ulcerative colitis  
 XX  
 SQ Sequence 22 BP; 6 A; 3 C; 7 G; 6 T; 0 U; 0 Other;  
 Query Match 100.0%; Score 22; DB 4; Length 22;  
 Best Local Similarity 100.0%; Pred. No. 0.24;  
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 OY 1 TGACTGTGAACGTTTCGAGATGA 22  
 DB 1 TGACTGTGAACGTTTCGAGATGA 22  
 Search completed: March 28, 2005, 22:51:34  
 Job time : 274 secs

GenCore version 5.1.6  
Copyright (c) 1993 - 2005 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: March 28, 2005, 22:38:52 ; Search time 94 Seconds  
(without alignments)  
382.958 Million cell updates/sec

Title: US-09-802-376-1

Perfect score: 22

Sequence: 1 tgactgtgaacgttcgagatga 22

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 1202784 seqs, 818138359 residues

Total number of hits satisfying chosen parameters: 2405568

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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- 3: /cgn2\_6/prodata1/ina/6A\_COMB.seq.\*
- 4: /cgn2\_6/prodata1/ina/6B\_COMB.seq.\*
- 5: /cgn2\_6/prodata1/ina/PCTUS\_COMB.seq.\*
- 6: /cgn2\_6/prodata1/ina/backfileseq.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	22	100.0	22	4	US-09-235-742-19
2	22	100.0	22	4	US-09-347-343-32
3	22	100.0	22	4	US-09-820-484-1
4	22	100.0	22	4	US-09-820-484-3
5	22	100.0	22	4	US-09-774-403A-1
6	22	100.0	22	4	US-09-296-477-2
7	22	100.0	22	4	US-09-308-036A-1
8	22	100.0	22	4	US-09-791-500-1
9	22	100.0	22	4	US-09-565-906-2
10	21	95.5	22	4	US-09-296-477-15
11	20.4	92.7	22	3	US-09-092-314-2
12	20.4	92.7	22	4	US-09-820-484-2
13	20.4	92.7	22	4	US-09-820-484-6
14	20.4	92.7	22	4	US-09-774-403A-2
15	20.4	92.7	22	4	US-09-296-477-1
16	20.4	92.7	22	4	US-09-296-477-5
17	20.4	92.7	22	4	US-09-296-477-6
18	20.4	92.7	22	4	US-09-791-500-4
19	20.4	92.7	22	4	US-09-791-500-5
20	20.4	92.7	22	4	US-09-791-500-6
21	20	90.9	22	4	US-09-296-477-16
22	19.4	88.2	22	4	US-09-296-477-12
23	18.8	85.5	22	3	US-09-092-314-1
24	18.8	85.5	22	3	US-09-092-314-3
25	18.8	85.5	22	3	US-09-092-314-10
26	18.8	85.5	22	4	US-09-235-742-20
27	18.8	85.5	22	4	US-09-347-343-33

28	18.8	85.5	22	4	US-09-820-484-7	Sequence 7, Appli
29	18.8	85.5	22	4	US-09-774-403A-3	Sequence 3, Appli
30	18.8	85.5	22	4	US-09-296-477-3	Sequence 3, Appli
31	18.8	85.5	22	4	US-09-296-477-8	Sequence 8, Appli
32	18.8	85.5	22	4	US-09-308-036A-2	Sequence 2, Appli
33	18.8	85.5	22	4	US-09-791-500-3	Sequence 3, Appli
34	18.8	85.5	22	4	US-09-791-500-8	Sequence 8, Appli
35	17.2	78.2	22	3	US-09-092-314-4	Sequence 4, Appli
36	17.2	78.2	22	4	US-09-296-477-9	Sequence 9, Appli
37	17.2	78.2	22	4	US-09-296-477-13	Sequence 13, Appli
38	17.2	78.2	22	4	US-09-791-500-9	Sequence 9, Appli
39	17.2	78.2	601	4	US-09-949-016-131826	Sequence 131826,
40	17.2	78.2	40493	4	US-09-949-016-15453	Sequence 15453, A
41	16.8	76.4	649	4	US-09-902-540-1286	Sequence 1286, Ap
42	16.2	73.6	601	4	US-09-949-016-204849	Sequence 204849, A
43	16.2	73.6	128723	4	US-09-949-016-17533	Sequence 17533, A
44	15.8	71.8	601	4	US-09-949-016-174829	Sequence 174829,
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ALIGNMENTS

RESULT 1

US-09-235-742-19

; Sequence 19, Application US/09235742

; Patent No. 6498148

; GENERAL INFORMATION:

; APPLICANT: Raz, Eval

; TITLE OF INVENTION: Immunization-Free Methods for Treating

; TITLE OF INVENTION: Antigen-Stimulated Inflammation in a Mammalian Host and

; TITLE OF INVENTION: Shifting the Host's Antigen Immune Responsiveness to a THI

; FILE REFERENCE: 6510-170CON4

; CURRENT APPLICATION NUMBER: US/09/235,742

; CURRENT FILING DATE: 1999-01-21

; EARLIER APPLICATION NUMBER: 08/927,120

; EARLIER FILING DATE: 1997-09-05

; EARLIER APPLICATION NUMBER: 08/593,554

; EARLIER FILING DATE: 1996-01-30

; EARLIER APPLICATION NUMBER: 08/725,968

; EARLIER FILING DATE: 1996-10-04

; EARLIER APPLICATION NUMBER: 60/028,118

; EARLIER FILING DATE: 1996-10-11

; NUMBER OF SEQ ID NOS: 20

; SOFTWARE: FastSeq for Windows Version 4.0

; SEQ ID NO 19

; LENGTH: 22

; TYPE: DNA

; ORGANISM: Artificial Sequence

; FEATURE:

; OTHER INFORMATION: Recombinant or Synthetic Sequence

US-09-235-742-19

Query Match 100.0%; Score 22; DB 4; Length 22;

Best Local Similarity 100.0%; Pred. No. 0.075;

Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTCCGAGATGA 22

Db 1 TGACTGTGAACGTTCCGAGATGA 22

RESULT 2

US-09-347-343-32

; Sequence 32, Application US/09347343A

; Patent No. 6514948

; GENERAL INFORMATION:

; APPLICANT: RAZ, Eval R.

; APPLICANT: KOBAYASHI, Hiroko

; TITLE OF INVENTION: METHOD FOR ENHANCING AN IMMUNE RESPONSE

; FILE REFERENCE: 30448.64US01

; CURRENT APPLICATION NUMBER: US/09/347,343A

; CURRENT FILING DATE: 1999-07-02  
; NUMBER OF SEQ ID NOS: 40  
; SOFTWARE: FastSeq for Windows Version 3.0  
; SEQ ID NO 32  
; LENGTH: 22  
; TYPE: DNA  
; ORGANISM: synthetic oligonucleotide  
US-09-347-343-32

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Best Local Similarity 100.0%; Pred. No. 0.075;  
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGAAGTGAACGTTTCGAGATGA 22  
Db 1 TGAAGTGAACGTTTCGAGATGA 22

RESULT 3  
US-09-820-484-1  
; Sequence 1, Application US/09820484  
; Patent No. 6534062  
; GENERAL INFORMATION:  
; APPLICANT: Raz, Eyal  
; APPLICANT: Cho, Hearn Jay  
; APPLICANT: Richman, Douglas  
; APPLICANT: Horner, Anthony A.  
; TITLE OF INVENTION: Method for Increasing a Cytotoxic T  
; FILE REFERENCE: 06510-188US1  
; CURRENT FILING DATE: 2001-03-28  
; PRIOR FILING DATE: 2001-03-28  
; PRIOR FILING DATE: 2000-03-28  
; PRIOR FILING DATE: 2000-03-28  
; PRIOR FILING DATE: 2000-05-11  
; PRIOR FILING DATE: 2000-05-11  
; PRIOR FILING DATE: 2000-07-05  
; NUMBER OF SEQ ID NOS: 8  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 1  
; LENGTH: 22  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Disulfide-linked phosphorothioate ISS-ODN  
; NAME/KEY: modified\_base  
; LOCATION: (1)...(1)  
; OTHER INFORMATION: disulfide thymine  
US-09-820-484-1

Query Match 100.0%; Score 22; DB 4; Length 22;  
Best Local Similarity 100.0%; Pred. No. 0.075;  
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGAAGTGAACGTTTCGAGATGA 22  
Db 1 TGAAGTGAACGTTTCGAGATGA 22

RESULT 4  
US-09-820-484-3  
; Sequence 3, Application US/09820484  
; Patent No. 6534062  
; GENERAL INFORMATION:  
; APPLICANT: Raz, Eyal  
; APPLICANT: Cho, Hearn Jay  
; APPLICANT: Richman, Douglas  
; APPLICANT: Horner, Anthony A.  
; TITLE OF INVENTION: Method for Increasing a Cytotoxic T  
; FILE REFERENCE: 06510-188US1  
; CURRENT FILING DATE: 2001-03-28  
; PRIOR FILING DATE: 2001-03-28  
; PRIOR FILING DATE: 2000-03-28  
; PRIOR FILING DATE: 2000-03-28  
; PRIOR FILING DATE: 2000-05-11  
; PRIOR FILING DATE: 2000-05-11  
; PRIOR FILING DATE: 2000-07-05  
; NUMBER OF SEQ ID NOS: 8  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 1  
; LENGTH: 22  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Disulfide-linked phosphorothioate ISS-ODN  
; NAME/KEY: modified\_base  
; LOCATION: (1)...(1)  
; OTHER INFORMATION: disulfide thymine  
US-09-820-484-1

; CURRENT FILING DATE: 2001-03-28  
; PRIOR FILING DATE: 2000-03-28  
; PRIOR FILING DATE: 2000-03-28  
; PRIOR FILING DATE: 2000-05-11  
; PRIOR FILING DATE: 2000-05-11  
; PRIOR FILING DATE: 2000-07-05  
; NUMBER OF SEQ ID NOS: 8  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 3  
; LENGTH: 22  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: phosphorothioate ISS-ODN  
US-09-820-484-3

Query Match 100.0%; Score 22; DB 4; Length 22;  
Best Local Similarity 100.0%; Pred. No. 0.075;  
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGAAGTGAACGTTTCGAGATGA 22  
Db 1 TGAAGTGAACGTTTCGAGATGA 22

RESULT 5  
US-09-774-403A-1  
; Sequence 1, Application US/09774403A  
; Patent No. 6552006  
; GENERAL INFORMATION:  
; APPLICANT: Eyal Raz  
; APPLICANT: Richard Kornbluth  
; APPLICANT: Antonio Catanzaro  
; APPLICANT: Tomoko Hayashi  
; APPLICANT: Dennis Carson  
; TITLE OF INVENTION: Immunomodulatory Polynucleotides in  
; FILE REFERENCE: UCAL166  
; CURRENT FILING DATE: 2002-04-15  
; PRIOR FILING DATE: 2000-01-31  
; PRIOR FILING DATE: 2000-01-31  
; NUMBER OF SEQ ID NOS: 7  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 1  
; LENGTH: 22  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Immunomodulatory sequence  
US-09-774-403A-1

Query Match 100.0%; Score 22; DB 4; Length 22;  
Best Local Similarity 100.0%; Pred. No. 0.075;  
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGAAGTGAACGTTTCGAGATGA 22  
Db 1 TGAAGTGAACGTTTCGAGATGA 22

RESULT 6  
US-09-296-477-2  
; Sequence 2, Application US/09296477A  
; Patent No. 6589940  
; GENERAL INFORMATION:  
; APPLICANT: RAZ, E.  
; APPLICANT: SCHWARTZ, D.  
; APPLICANT: ROMAN, M.  
; APPLICANT: DINA, D.  
; TITLE OF INVENTION: IMMUNOSTIMULATORY OLIGONUCLEOTIDES,  
; COMPOSITIONS THEREOF AND METHODS OF USE

```
; TITLE OF INVENTION: THEREOF
; FILE REFERENCE: 37782000420
; CURRENT APPLICATION NUMBER: US/09/296,477A
; CURRENT FILING DATE: 1999-04-22
; EARLIER APPLICATION NUMBER: 09/092,329
; EARLIER FILING DATE: 1998-06-05
; EARLIER APPLICATION NUMBER: 60/048,793
; EARLIER FILING DATE: 1997-06-06
; NUMBER OF SEQ ID NOS: 21
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 2
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic construct
US-09-296-477-2

Query Match      100.0%; Score 22; DB 4; Length 22;
Best Local Similarity 100.0%; Pred. No. 0.075;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 TGA CTGTGAACGTTTCGAGATGA 22
Db      1 TGA CTGTGAACGTTTCGAGATGA 22

RESULT 7
US-09-308-036A-1
; Sequence 1, Application US/09308036A
; Patent No. 6610661
; GENERAL INFORMATION:
; APPLICANT: Carson, Dennis A.
; APPLICANT: Raz, Eyal
; APPLICANT: Roman, Mark
; TITLE OF INVENTION: Polynucleotide/Immunomodulatory Molecule Conjugates
; FILE REFERENCE: 6510-172CIP
; CURRENT APPLICATION NUMBER: US/09/308,036A
; CURRENT FILING DATE: 2000-02-16
; PRIOR APPLICATION NUMBER: PCT/US97/19004
; PRIOR FILING DATE: 1997-10-09
; PRIOR APPLICATION NUMBER: 60/028,118
; PRIOR FILING DATE: 1996-10-11
; NUMBER OF SEQ ID NOS: 2
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: DY1018 polynucleotide
US-09-308-036A-1

Query Match      100.0%; Score 22; DB 4; Length 22;
Best Local Similarity 100.0%; Pred. No. 0.075;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 TGA CTGTGAACGTTTCGAGATGA 22
Db      1 TGA CTGTGAACGTTTCGAGATGA 22

RESULT 8
US-09-791-500-1
; Sequence 1, Application US/09791500
; Patent No. 6613751
; GENERAL INFORMATION:
; APPLICANT: Rachmilewitz, Daniel
; APPLICANT: Raz, Eyal
; TITLE OF INVENTION: Method for Treating Inflammatory Bowel
; TITLE OF INVENTION: Disease and Other Forms of Gastrointestinal Inflammation.
; FILE REFERENCE: 6510-202US1

; TITLE OF INVENTION: THEREOF
; FILE REFERENCE: 37782000420
; CURRENT APPLICATION NUMBER: US/09/296,477A
; CURRENT FILING DATE: 1999-04-22
; EARLIER APPLICATION NUMBER: 09/092,329
; EARLIER FILING DATE: 1998-06-05
; EARLIER APPLICATION NUMBER: 60/048,793
; EARLIER FILING DATE: 1997-06-06
; NUMBER OF SEQ ID NOS: 21
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 2
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic polynucleotide sequence
US-09-791-500-1

Query Match      100.0%; Score 22; DB 4; Length 22;
Best Local Similarity 100.0%; Pred. No. 0.075;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 TGA CTGTGAACGTTTCGAGATGA 22
Db      1 TGA CTGTGAACGTTTCGAGATGA 22

RESULT 9
US-09-565-906-2
; Sequence 2, Application US/09565906
; Patent No. 6737066
; GENERAL INFORMATION:
; APPLICANT: Moss, Ronald B.
; TITLE OF INVENTION: HIV Immunogenic Compositions and Methods
; FILE REFERENCE: P-IM 4029
; CURRENT APPLICATION NUMBER: US/09/565,906
; CURRENT FILING DATE: 2000-05-05
; PRIOR APPLICATION NUMBER: US 60/132,762
; PRIOR FILING DATE: 1999-05-06
; PRIOR APPLICATION NUMBER: US 60/150,667
; PRIOR FILING DATE: 1999-08-25
; NUMBER OF SEQ ID NOS: 4
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: phosphorothioate-modified synthetic
; OTHER INFORMATION: oligodeoxynucleotide
US-09-565-906-2

Query Match      100.0%; Score 22; DB 4; Length 22;
Best Local Similarity 100.0%; Pred. No. 0.075;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 TGA CTGTGAACGTTTCGAGATGA 22
Db      1 TGA CTGTGAACGTTTCGAGATGA 22

RESULT 10
US-09-296-477-15
; Sequence 15, Application US/09296477A
; Patent No. 6589940
; GENERAL INFORMATION:
; APPLICANT: RAZ, E.
; APPLICANT: SCHWARTZ, D.
; APPLICANT: ROMAN, M.
; APPLICANT: DINA, D.
; TITLE OF INVENTION: IMMUNOSTIMULATORY OLIGONUCLEOTIDES,
; TITLE OF INVENTION: COMPOSITIONS THEREOF AND METHODS OF USE
; TITLE OF INVENTION: THEREOF
; FILE REFERENCE: 37782000420
; CURRENT APPLICATION NUMBER: US/09/296,477A
; CURRENT FILING DATE: 1999-04-22
; EARLIER APPLICATION NUMBER: 09/092,329
; EARLIER FILING DATE: 1998-06-05
; EARLIER APPLICATION NUMBER: 60/048,793
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; EARLIER FILING DATE: 1997-06-06
; NUMBER OF SEQ ID NOS: 21
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 15
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic construct
; NAME/KEY: modified base
; LOCATION: (11)...(11)
; OTHER INFORMATION: 5-bromocytosine
US-09-296-477-15

Query Match          95.5%; Score 21; DB 4; Length 22;
Best Local Similarity 95.5%; Pred. No. 0.24;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTTCGAGATGA 22
   ||||| ||||| ||||| |||||
DB 1 TGACTGTGAAGTTTCGAGATGA 22

RESULT 11
US-09-092-314-2
; Sequence 2, Application US/09092314
; Patent No. 6225292
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; APPLICANT: Roman, Mark
; TITLE OF INVENTION: Inhibitors of DNA Immunostimulatory
; TITLE OF INVENTION: Sequence Activity
; Patent No. 6225292
; FILE REFERENCE: 6510-173US1
; CURRENT APPLICATION NUMBER: US/09/092,314
; CURRENT FILING DATE: 1998-06-05
; PRIOR APPLICATION NUMBER: 60/048,794
; PRIOR FILING DATE: 1997-06-06
; NUMBER OF SEQ ID NOS: 11
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide
US-09-092-314-2

Query Match          92.7%; Score 20.4; DB 3; Length 22;
Best Local Similarity 95.5%; Pred. No. 0.49;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTTCGAGATGA 22
   ||||| ||||| ||||| |||||
DB 1 TGACTGTGAACGTTAGAGATGA 22

RESULT 12
US-09-820-484-2
; Sequence 2, Application US/09820484
; Patent No. 6534062
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; APPLICANT: Cho, Hearn Jay
; APPLICANT: Richman, Douglas
; APPLICANT: Horner, Anthony A.
; TITLE OF INVENTION: Method for Increasing a Cytotoxic T
; TITLE OF INVENTION: Lymphocyte Response in vivo.
; FILE REFERENCE: 06510-188US1
; CURRENT APPLICATION NUMBER: US/09/820,484
; CURRENT FILING DATE: 2001-03-28
; PRIOR APPLICATION NUMBER: US/09/820,484
; PRIOR FILING DATE: 2000-07-05
; NUMBER OF SEQ ID NOS: 8
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 6
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: mutated control ODN
US-09-820-484-2

Query Match          92.7%; Score 20.4; DB 4; Length 22;
Best Local Similarity 95.5%; Pred. No. 0.49;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTTCGAGATGA 22
   ||||| ||||| ||||| |||||
DB 1 TGACTGTGAACCTTCGAGATGA 22

RESULT 14
US-09-774-403A-2
; Sequence 2, Application US/09774403A
; Patent No. 6552006
; GENERAL INFORMATION:
; APPLICANT: Eyal Raz
; APPLICANT: Richard Kornbluth

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; EARLIER FILING DATE: 1997-06-06
; NUMBER OF SEQ ID NOS: 21
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 15
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic construct
; NAME/KEY: modified base
; LOCATION: (11)...(11)
; OTHER INFORMATION: 5-bromocytosine
US-09-296-477-15

Query Match          95.5%; Score 21; DB 4; Length 22;
Best Local Similarity 95.5%; Pred. No. 0.24;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTTCGAGATGA 22
   ||||| ||||| ||||| |||||
DB 1 TGACTGTGAAGTTTCGAGATGA 22

RESULT 11
US-09-092-314-2
; Sequence 2, Application US/09092314
; Patent No. 6225292
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; APPLICANT: Roman, Mark
; TITLE OF INVENTION: Inhibitors of DNA Immunostimulatory
; TITLE OF INVENTION: Sequence Activity
; Patent No. 6225292
; FILE REFERENCE: 6510-173US1
; CURRENT APPLICATION NUMBER: US/09/092,314
; CURRENT FILING DATE: 1998-06-05
; PRIOR APPLICATION NUMBER: 60/048,794
; PRIOR FILING DATE: 1997-06-06
; NUMBER OF SEQ ID NOS: 11
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide
US-09-092-314-2

Query Match          92.7%; Score 20.4; DB 3; Length 22;
Best Local Similarity 95.5%; Pred. No. 0.49;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTTCGAGATGA 22
   ||||| ||||| ||||| |||||
DB 1 TGACTGTGAACGTTAGAGATGA 22

RESULT 12
US-09-820-484-2
; Sequence 2, Application US/09820484
; Patent No. 6534062
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; APPLICANT: Cho, Hearn Jay
; APPLICANT: Richman, Douglas
; APPLICANT: Horner, Anthony A.
; TITLE OF INVENTION: Method for Increasing a Cytotoxic T
; TITLE OF INVENTION: Lymphocyte Response in vivo.
; FILE REFERENCE: 06510-188US1
; CURRENT APPLICATION NUMBER: US/09/820,484
; CURRENT FILING DATE: 2001-03-28
; PRIOR APPLICATION NUMBER: US/09/820,484
; PRIOR FILING DATE: 2000-07-05
; NUMBER OF SEQ ID NOS: 8
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 6
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: mutated control ODN
US-09-820-484-2

Query Match          92.7%; Score 20.4; DB 4; Length 22;
Best Local Similarity 95.5%; Pred. No. 0.49;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTTCGAGATGA 22
   ||||| ||||| ||||| |||||
DB 1 TGACTGTGAACCTTCGAGATGA 22

RESULT 13
US-09-820-484-6
; Sequence 6, Application US/09820484
; Patent No. 6534062
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; APPLICANT: Cho, Hearn Jay
; APPLICANT: Richman, Douglas
; APPLICANT: Horner, Anthony A.
; TITLE OF INVENTION: Method for Increasing a Cytotoxic T
; TITLE OF INVENTION: Lymphocyte Response in vivo.
; FILE REFERENCE: 06510-188US1
; CURRENT APPLICATION NUMBER: US/09/820,484
; CURRENT FILING DATE: 2001-03-28
; PRIOR APPLICATION NUMBER: US/09/820,484
; PRIOR FILING DATE: 2000-03-28
; PRIOR APPLICATION NUMBER: US/09/820,484
; PRIOR FILING DATE: 2000-05-11
; PRIOR APPLICATION NUMBER: US/09/820,484
; PRIOR FILING DATE: 2000-07-05
; NUMBER OF SEQ ID NOS: 8
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 6
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: mutated control ODN
US-09-820-484-6

Query Match          92.7%; Score 20.4; DB 4; Length 22;
Best Local Similarity 95.5%; Pred. No. 0.49;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTTCGAGATGA 22
   ||||| ||||| ||||| |||||
DB 1 TGACTGTGAACCTTCGAGATGA 22

RESULT 14
US-09-774-403A-2
; Sequence 2, Application US/09774403A
; Patent No. 6552006
; GENERAL INFORMATION:
; APPLICANT: Eyal Raz
; APPLICANT: Richard Kornbluth

```

APPLICANT: Antonio Catanzaro  
APPLICANT: Tomoko Hayashi  
APPLICANT: Dennis Carson  
TITLE OF INVENTION: Immunomodulatory Polynucleotides in  
FILE REFERENCE: UCAL166  
CURRENT APPLICATION NUMBER: US/09/774,403A  
CURRENT FILING DATE: 2002-04-15  
PRIOR APPLICATION NUMBER: 60/179,353  
PRIOR FILING DATE: 2000-01-31  
NUMBER OF SEQ ID NOS: 7  
SOFTWARE: FastSeq for Windows Version 4.0  
SEQ ID NO 2  
LENGTH: 22  
TYPE: DNA  
ORGANISM: Artificial Sequence  
FEATURE:  
OTHER INFORMATION: Control sequence  
US-09-774-403A-2

Query Match 92.7%; Score 20.4; DB 4; Length 22;  
Best Local Similarity 95.5%; Pred. No. 0.49;  
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTTCGAGATGA 22  
DB 1 TGACTGTGAAGGTTTCGAGATGA 22

RESULT 15  
US-09-296-477-1  
Sequence 1, Application US/09296477A  
Patent No. 6589940  
GENERAL INFORMATION:  
APPLICANT: RAZ, E.  
APPLICANT: SCHWARTZ, D.  
APPLICANT: ROMAN, M.  
APPLICANT: DINA, D.  
TITLE OF INVENTION: IMMUNOSTIMULATORY OLIGONUCLEOTIDES,  
TITLE OF INVENTION: COMPOSITIONS THEREOF AND METHODS OF USE  
FILE REFERENCE: 37788200420  
CURRENT APPLICATION NUMBER: US/09/296,477A  
CURRENT FILING DATE: 1999-04-22  
EARLIER APPLICATION NUMBER: 09/092,329  
EARLIER FILING DATE: 1998-06-05  
EARLIER APPLICATION NUMBER: 60/048,793  
EARLIER FILING DATE: 1997-06-06  
NUMBER OF SEQ ID NOS: 21  
SOFTWARE: FastSeq for Windows Version 3.0  
SEQ ID NO 1  
LENGTH: 22  
TYPE: DNA  
ORGANISM: Artificial Sequence  
FEATURE:  
OTHER INFORMATION: Synthetic construct  
US-09-296-477-1

Query Match 92.7%; Score 20.4; DB 4; Length 22;  
Best Local Similarity 95.5%; Pred. No. 0.49;  
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTTCGAGATGA 22  
DB 1 TGACCGTGAACGTTTCGAGATGA 22

Search completed: March 28, 2005, 23:51:57  
Job time : 95 secs

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GenCore version 5.1.6  
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OM nucleic - nucleic search, using sw model

Run on: March 28, 2005, 22:47:02 ; Search time 319 Seconds  
(without alignments)  
410.988 Million cell updates/sec

Title: US-09-802-376-1

Perfect score: 22

Sequence: 1 TGACTGTGAACGTCGAGATGA 22

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 5552208 seqs, 2979665951 residues

Total number of hits satisfying chosen parameters: 11104416

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications NA:\*

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17: /cgn2\_6/ptodata/2/pubpna/US10E\_PUBCOMB.seq:  
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

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2	22	100.0	22	9	US-09-802-685-1
3	22	100.0	22	9	US-09-791-500-1
4	22	100.0	22	9	US-09-802-376-1
5	22	100.0	22	9	US-09-774-403A-1
6	22	100.0	22	9	US-09-802-370-1
7	22	100.0	22	9	US-09-802-445-1
8	22	100.0	22	9	US-09-820-484-1
9	22	100.0	22	9	US-09-820-484-3
10	22	100.0	22	9	US-09-828-505-1
11	22	100.0	22	9	US-09-967-881-2

12	22	100.0	22	10	US-09-927-422A-1	Sequence 1, Appli
13	22	100.0	22	10	US-09-738-046A-3	Sequence 3, Appli
14	22	100.0	22	10	US-09-927-884-1	Sequence 1, Appli
15	22	100.0	22	10	US-09-802-359-1	Sequence 1, Appli
16	22	100.0	22	10	US-09-967-464-19	Sequence 19, Appli
17	22	100.0	22	10	US-09-848-986-1	Sequence 1, Appli
18	22	100.0	22	14	US-10-056-420-4	Sequence 4, Appli
19	22	100.0	22	14	US-10-033-243-2	Sequence 2, Appli
20	22	100.0	22	14	US-10-033-243-40	Sequence 40, Appli
21	22	100.0	22	14	US-10-033-243-59	Sequence 59, Appli
22	22	100.0	22	14	US-10-214-288-1	Sequence 1, Appli
23	22	100.0	22	14	US-10-099-512-1	Sequence 1, Appli
24	22	100.0	22	14	US-10-229-208-19	Sequence 19, Appli
25	22	100.0	22	15	US-10-233-117-32	Sequence 32, Appli
26	22	100.0	22	15	US-10-233-121A-1	Sequence 1, Appli
27	22	100.0	22	15	US-10-219-143-1	Sequence 2, Appli
28	22	100.0	22	15	US-10-214-799-2	Sequence 1, Appli
29	22	100.0	22	15	US-10-340-275-1	Sequence 1, Appli
30	22	100.0	22	15	US-10-340-275-3	Sequence 3, Appli
31	22	100.0	22	15	US-10-333-885-1	Sequence 1, Appli
32	22	100.0	22	15	US-10-339-885-3	Sequence 3, Appli
33	22	100.0	22	16	US-10-176-883-2	Sequence 2, Appli
34	22	100.0	22	16	US-10-176-883-24	Sequence 24, Appli
35	22	100.0	22	16	US-10-176-883-79	Sequence 79, Appli
36	22	100.0	22	16	US-10-176-883-134	Sequence 134, App
37	22	100.0	22	16	US-10-412-151-1	Sequence 1, Appli
38	22	100.0	22	16	US-10-177-826-2	Sequence 2, Appli
39	22	100.0	22	16	US-10-177-826-24	Sequence 24, Appli
40	22	100.0	22	16	US-10-177-826-79	Sequence 79, Appli
41	22	100.0	22	16	US-10-177-826-134	Sequence 134, App
42	22	100.0	22	17	US-10-353-917-1	Sequence 1, Appli
43	22	100.0	22	17	US-10-357-760-1	Sequence 1, Appli
44	22	100.0	22	17	US-10-328-578-2	Sequence 2, Appli
45	22	100.0	22	17	US-10-328-578-24	Sequence 24, Appli

#### ALIGNMENTS

#### RESULT 1

US-09-802-686-1  
; Sequence 1, Application US/09802686  
; Patent No. US200010046967A1  
; GENERAL INFORMATION:  
; APPLICANT: Dynavax Technologies Corporation  
; APPLICANT: Van Nest, Gary  
; TITLE OF INVENTION: METHODS OF PREVENTING AND TREATING  
; TITLE OF INVENTION: RESPIRATORY VIRAL INFECTION USING IMMUNOMODULATORY  
; TITLE OF INVENTION: POLYNUCLEOTIDE SEQUENCES  
; FILE REFERENCE: 377882000900  
; CURRENT APPLICATION NUMBER: US/09/802,686  
; CURRENT FILING DATE: 2001-03-09  
; PRIOR APPLICATION NUMBER: 60/188,583  
; PRIOR FILING DATE: 2000-03-10  
; NUMBER OF SEQ ID NOS: 10  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 1  
; LENGTH: 22  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Polynucleotide containing CG  
US-09-802-686-1

Query Match 100.0%; Score 22; DB 9; Length 22;

Best Local Similarity 100.0%; Pred. No. 0.4;

Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTCGAGATGA 22

Db 1 TGACTGTGAACGTCGAGATGA 22

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RESULT 2
US-09-802-685-1
; CURRENT APPLICATION NUMBER: US/09802685
; CURRENT FILING DATE: 2001-03-09
; PRIOR APPLICATION NUMBER: 60/188,557
; PRIOR FILING DATE: 2000-03-10
; NUMBER OF SEQ ID NOS: 11
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Polynucleotide containing CG
US-09-802-685-1
Query Match      100.0%; Score 22; DB 9; Length 22;
Best Local Similarity 100.0%; Pred. No. 0.4;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGAAGTGTGAACGTTTCGAGATGA 22
   |||||
Db 1 TGAAGTGTGAACGTTTCGAGATGA 22

RESULT 3
US-09-791-500-1
; Sequence 1, Application US/09791500
; Patent No. US20020042387A1
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; APPLICANT: Rachmilewitz, Daniel
; TITLE OF INVENTION: Method for Treating Inflammatory Bowel
; TITLE OF INVENTION: Disease and Other Forms of Gastrointestinal Inflammation.
; FILE REFERENCE: 6510-202US1
; CURRENT APPLICATION NUMBER: US/09/791,500
; CURRENT FILING DATE: 2001-02-22
; NUMBER OF SEQ ID NOS: 39
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: synthetic polynucleotide sequence
US-09-791-500-1
Query Match      100.0%; Score 22; DB 9; Length 22;
Best Local Similarity 100.0%; Pred. No. 0.4;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGAAGTGTGAACGTTTCGAGATGA 22
   |||||
Db 1 TGAAGTGTGAACGTTTCGAGATGA 22

RESULT 4
US-09-802-376-1
; Sequence 1, Application US/09802376
; Patent No. US20020055477A1
; GENERAL INFORMATION:
; APPLICANT: Van Nest, Gary
; APPLICANT: Tuck, Stephen
; TITLE OF INVENTION: IMMUNOMODULATORY FORMULATIONS AND METHODS FOR USE THEREOF
; FILE REFERENCE: 37788201700
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; CURRENT APPLICATION NUMBER: US/09/802,376
; CURRENT FILING DATE: 2001-03-09
; PRIOR APPLICATION NUMBER: 60/188,557
; PRIOR FILING DATE: 2000-03-10
; NUMBER OF SEQ ID NOS: 11
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Polynucleotide containing CG
US-09-802-376-1
Query Match      100.0%; Score 22; DB 9; Length 22;
Best Local Similarity 100.0%; Pred. No. 0.4;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGAAGTGTGAACGTTTCGAGATGA 22
   |||||
Db 1 TGAAGTGTGAACGTTTCGAGATGA 22

RESULT 5
US-09-774-403A-1
; Sequence 1, Application US/09774403A
; Publication No. US20020086295A1
; GENERAL INFORMATION:
; APPLICANT: Eyal Raz
; APPLICANT: Richard Kornbluth
; APPLICANT: Antonio Catanzaro
; APPLICANT: Tomoko Hayashi
; APPLICANT: Dennis Carson
; TITLE OF INVENTION: Immunomodulatory Polynucleotides in
; TITLE OF INVENTION: Treatment of Infection by an Intracellular Pathogen
; FILE REFERENCE: UCAL166
; CURRENT APPLICATION NUMBER: US/09/774,403A
; CURRENT FILING DATE: 2002-04-15
; PRIOR APPLICATION NUMBER: 60/179,353
; PRIOR FILING DATE: 2000-01-31
; NUMBER OF SEQ ID NOS: 7
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Immunomodulatory sequence
US-09-774-403A-1
Query Match      100.0%; Score 22; DB 9; Length 22;
Best Local Similarity 100.0%; Pred. No. 0.4;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGAAGTGTGAACGTTTCGAGATGA 22
   |||||
Db 1 TGAAGTGTGAACGTTTCGAGATGA 22

RESULT 6
US-09-802-370-1
; Sequence 1, Application US/09802370
; Patent No. US20020098199A1
; GENERAL INFORMATION:
; APPLICANT: Van Nest, Gary
; APPLICANT: Biden, Joseph J. Jr.
; TITLE OF INVENTION: METHODS OF SUPPRESSING HEPATITIS VIRUS
; TITLE OF INVENTION: INFECTION USING IMMUNOMODULATORY POLYNUCLEOTIDE SEQUENCES
; FILE REFERENCE: 377882001200
; CURRENT APPLICATION NUMBER: US/09/802,370
; CURRENT FILING DATE: 2001-09-24
; PRIOR APPLICATION NUMBER: 60/188,301
; PRIOR FILING DATE: 2000-03-10
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; NUMBER OF SEQ ID NOS: 8  
; SOFTWARE: FastSEQ for Windows Version 4.0  
; SEQ ID NO 1  
; LENGTH: 22  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Polynucleotide containing CG  
US-09-802-370-1

Query Match 100.0%; Score 22; DB 9; Length 22;  
Best Local Similarity 100.0%; Pred. No. 0.4;  
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTTCGAGATGA 22  
|||  
Db 1 TGACTGTGAACGTTTCGAGATGA 22

## RESULT 7

US-09-802-445-1  
; Sequence 1, Application US/09802445  
; Patent No. US20020107212A1  
; GENERAL INFORMATION:  
; APPLICANT: Van Nest, Gary  
; APPLICANT: Eiden, Joseph J. Jr.  
; TITLE OF INVENTION: METHODS OF REDUCING PAPILLOMAVIRUS INFECTION USING IMMUNOMODULATORS  
; FILE REFERENCE: 37782001300  
; CURRENT APPLICATION NUMBER: US/09/802,445  
; CURRENT FILING DATE: 2001-09-24  
; PRIOR APPLICATION NUMBER: 60/188,265  
; PRIOR FILING DATE: 2000-03-10  
; NUMBER OF SEQ ID NOS: 8  
; SOFTWARE: FastSEQ for Windows Version 4.0  
; SEQ ID NO 1  
; LENGTH: 22  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Polynucleotide containing CG  
US-09-802-445-1

Query Match 100.0%; Score 22; DB 9; Length 22;  
Best Local Similarity 100.0%; Pred. No. 0.4;  
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTTCGAGATGA 22  
|||  
Db 1 TGACTGTGAACGTTTCGAGATGA 22

## RESULT 8

US-09-820-484-1  
; Sequence 1, Application US/09820484  
; Patent No. US20020142977A1  
; GENERAL INFORMATION:  
; APPLICANT: Raz, Eyal  
; APPLICANT: Cho, Hearn Jay  
; APPLICANT: Richman, Douglas  
; APPLICANT: Horner, Anthony A.  
; TITLE OF INVENTION: Method for Increasing a Cytotoxic T  
; TITLE OF INVENTION: Lymphocyte Response in vivo.  
; FILE REFERENCE: 06510-188US1  
; CURRENT APPLICATION NUMBER: US/09/820,484  
; CURRENT FILING DATE: 2001-03-28  
; PRIOR APPLICATION NUMBER: US 60/192,537  
; PRIOR FILING DATE: 2000-03-28  
; PRIOR APPLICATION NUMBER: US 60/203,567  
; PRIOR FILING DATE: 2000-05-11  
; PRIOR APPLICATION NUMBER: US 60/215,895  
; PRIOR FILING DATE: 2000-07-05  
; NUMBER OF SEQ ID NOS: 8

; SOFTWARE: FastSEQ for Windows Version 4.0  
; SEQ ID NO 1  
; LENGTH: 22  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Disulfide-linked phosphorothioate ISS-ODN  
; NAME/KEY: modified base  
; LOCATION: (1)...(1)  
; OTHER INFORMATION: disulfide thymine  
US-09-820-484-1

Query Match 100.0%; Score 22; DB 9; Length 22;  
Best Local Similarity 100.0%; Pred. No. 0.4;  
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTTCGAGATGA 22  
|||  
Db 1 TGACTGTGAACGTTTCGAGATGA 22

## RESULT 9

US-09-820-484-3  
; Sequence 3, Application US/09820484  
; Patent No. US20020142977A1  
; GENERAL INFORMATION:  
; APPLICANT: Raz, Eyal  
; APPLICANT: Cho, Hearn Jay  
; APPLICANT: Richman, Douglas  
; APPLICANT: Horner, Anthony A.  
; TITLE OF INVENTION: Method for Increasing a Cytotoxic T  
; TITLE OF INVENTION: Lymphocyte Response in vivo.  
; FILE REFERENCE: 06510-188US1  
; CURRENT APPLICATION NUMBER: US/09/820,484  
; CURRENT FILING DATE: 2001-03-28  
; PRIOR APPLICATION NUMBER: US 60/192,537  
; PRIOR FILING DATE: 2000-03-28  
; PRIOR APPLICATION NUMBER: US 60/203,567  
; PRIOR FILING DATE: 2000-05-11  
; PRIOR APPLICATION NUMBER: US 60/215,895  
; PRIOR FILING DATE: 2000-07-05  
; NUMBER OF SEQ ID NOS: 8  
; SOFTWARE: FastSEQ for Windows Version 4.0  
; SEQ ID NO 3  
; LENGTH: 22  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: phosphorothioate ISS-ODN  
US-09-820-484-3

Query Match 100.0%; Score 22; DB 9; Length 22;  
Best Local Similarity 100.0%; Pred. No. 0.4;  
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTTCGAGATGA 22  
|||  
Db 1 TGACTGTGAACGTTTCGAGATGA 22

## RESULT 10

US-09-828-505-1  
; Sequence 1, Application US/09828505  
; Patent No. US20020142978A1  
; GENERAL INFORMATION:  
; APPLICANT: Raz, Eyal  
; APPLICANT: Takabayashi, Kenji  
; APPLICANT: Nguyen, Minh-Duc  
; TITLE OF INVENTION: Synergistic Improvements to  
; TITLE OF INVENTION: Polynucleotide Vaccines  
; FILE REFERENCE: 6510-203  
; CURRENT APPLICATION NUMBER: US/09/828,505  
; CURRENT FILING DATE: 2001-04-06

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; PRIOR APPLICATION NUMBER: 60/195,890
; PRIOR FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 4
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Immunomodulatory nucleic acid sequence
US-09-828-505-1

Query Match      100.0%; Score 22; DB 9; Length 22;
Best Local Similarity 100.0%; Pred. No. 0.4;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGAAGTGAACGTTTCGAGATGA 22
Db 1 TGAAGTGAACGTTTCGAGATGA 22

RESULT 11
US-09-967-881-2
; Sequence 2, Application US/09967881
; Publication No. US20020192184A1
; GENERAL INFORMATION:
; APPLICANT: Assistance Publique - Hopitaux de Paris
; APPLICANT: Institut National de la Sante et de la Recherche M
; APPLICANT: Carpentier, Ancoine
; TITLE OF INVENTION: Use of Stabilised Oligonucleotides for Preparing A Medicament wit
; TITLE OF INVENTION: Antitumor Activity
; FILE REFERENCE: 267/246 US
; CURRENT APPLICATION NUMBER: US/09/967,881
; CURRENT FILING DATE: 2001-09-28
; NUMBER OF SEQ ID NOS: 48
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 2
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Oligodeoxynucleotide
US-09-967-881-2

Query Match      100.0%; Score 22; DB 9; Length 22;
Best Local Similarity 100.0%; Pred. No. 0.4;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGAAGTGAACGTTTCGAGATGA 22
Db 1 TGAAGTGAACGTTTCGAGATGA 22

RESULT 12
US-09-927-422A-1
; Sequence 1, Application US/0927422A
; Publication No. US2003002852A1
; GENERAL INFORMATION:
; APPLICANT: Van Nest, Gary
; APPLICANT: Tuck, Stephen
; APPLICANT: Fearon, Karen L.
; APPLICANT: Dina, Dino
; TITLE OF INVENTION: BIODEGRADABLE IMMUNOMODULATORY
; TITLE OF INVENTION: FORMULATIONS AND METHODS FOR USE THEREOF
; FILE REFERENCE: 377882001420
; CURRENT APPLICATION NUMBER: US/09/927,422A
; CURRENT FILING DATE: 2001-08-10
; PRIOR APPLICATION NUMBER: U.S. 09/802,359
; PRIOR FILING DATE: 2001-03-09
; PRIOR APPLICATION NUMBER: U.S. 60/188,30
; PRIOR FILING DATE: 2000-03-10
; NUMBER OF SEQ ID NOS: 23
; SOFTWARE: FastSEQ for Windows Version 4.0

; SEQ ID NO 1
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Polynucleotide containing CG
US-09-927-422A-1

Query Match      100.0%; Score 22; DB 10; Length 22;
Best Local Similarity 100.0%; Pred. No. 0.4;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGAAGTGAACGTTTCGAGATGA 22
Db 1 TGAAGTGAACGTTTCGAGATGA 22

RESULT 13
US-09-738-046A-3
; Sequence 3, Application US/09738046A
; Publication No. US20030054007A1
; GENERAL INFORMATION:
; APPLICANT: FELGNER, PHILIP L.
; APPLICANT: ZELPHATI, OLIVIER
; TITLE OF INVENTION: INTRACELLULAR PROTEIN DELIVERY
; TITLE OF INVENTION: COMPOSITIONS AND METHODS OF USE
; FILE REFERENCE: GTSYS.004A
; CURRENT APPLICATION NUMBER: US/09/738,046A
; CURRENT FILING DATE: 2000-12-15
; NUMBER OF SEQ ID NOS: 3
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: artificial sequence containing CpG sequence
US-09-738-046A-3

Query Match      100.0%; Score 22; DB 10; Length 22;
Best Local Similarity 100.0%; Pred. No. 0.4;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGAAGTGAACGTTTCGAGATGA 22
Db 1 TGAAGTGAACGTTTCGAGATGA 22

RESULT 14
US-09-927-884-1
; Sequence 1, Application US/09927884
; Publication No. US20030059773A1
; GENERAL INFORMATION:
; APPLICANT: Van Nest, Gary
; APPLICANT: Tuck, Stephen
; APPLICANT: Fearon, Karen L.
; APPLICANT: Dina, Dino
; TITLE OF INVENTION: IMMUNOMODULATORY FORMULATIONS AND
; TITLE OF INVENTION: METHODS FOR USE THEREOF
; FILE REFERENCE: 377882001720
; CURRENT APPLICATION NUMBER: US/09/927,884
; CURRENT FILING DATE: 2001-08-10
; PRIOR APPLICATION NUMBER: U.S. 09/802,376
; PRIOR FILING DATE: 2001-03-09
; PRIOR APPLICATION NUMBER: U.S. 60/188,557
; PRIOR FILING DATE: 2000-03-10
; NUMBER OF SEQ ID NOS: 14
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
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; OTHER INFORMATION: Polynucleotide containing CG
US-09-927-884-1
Query Match      100.0%; Score 22; DB 10; Length 22;
Best Local Similarity 100.0%; Pred. No. 0.4;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 TGA CTGTG AAG GTTCG AGATGA 22
      |||||
Db      1 TGA CTGTG AAG GTTCG AGATGA 22
      |||||

RESULT 15
US-09-802-359-1
; Sequence 1, Application US/09802359
; Publication No. US20030129251A1
; GENERAL INFORMATION:
; APPLICANT: Van Nest, Gary
; APPLICANT: Tuck, Stephen
; TITLE OF INVENTION: IMMUNOMODULATORY FORMULATIONS AND METHODS FOR USE THEREOF
; FILE REFERENCE: 37788201400
; CURRENT APPLICATION NUMBER: US/09/802,359
; CURRENT FILING DATE: 2001-03-09
; PRIOR APPLICATION NUMBER: 60/188,303
; PRIOR FILING DATE: 2000-03-10
; NUMBER OF SEQ ID NOS: 11
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Polynucleotide containing CG
US-09-802-359-1

Query Match      100.0%; Score 22; DB 10; Length 22;
Best Local Similarity 100.0%; Pred. No. 0.4;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 TGA CTGTG AAG GTTCG AGATGA 22
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Db      1 TGA CTGTG AAG GTTCG AGATGA 22
      |||||

Search completed: March 28, 2005, 23:57:24
Job time : 321 secs
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GenCore version 5.1.6  
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: March 28, 2005, 22:36:17 ; Search time 1991 Seconds  
(without alignments)  
420.600 Million cell updates/sec

Title: US-09-802-376-1

Perfect score: 22  
Sequence: 1 tgactgtgaacgttcgagatga 22

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 69479088

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : EST.\*

1: gb\_est1.\*  
2: gb\_est2.\*  
3: gb\_hic.\*  
4: gb\_est3.\*  
5: gb\_est4.\*  
6: gb\_est5.\*  
7: gb\_est6.\*  
8: gb\_gss1.\*  
9: gb\_gss2.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	18.8	85.5	521	8	BH859011 S5_182b
2	18.4	83.6	571	4	BM042508 603615795
3	17.8	80.9	492	9	CE751403 tigr-gss-
4	17.8	80.9	561	8	AZ755668 ev02009.x
5	17.8	80.9	867	8	BZ558601 pa98401.2
6	17.8	80.9	961	4	BF971856 602240444
7	17.4	79.1	489	8	AZ060178 RFCI-23-4
8	17.4	79.1	530	8	AZ886419 RFCI-23-1
9	17.2	78.2	374	8	AQ245026 HS_2056 B
10	17.2	78.2	408	8	AZ536502
11	17.2	78.2	424	2	BE723539
12	17.2	78.2	427	7	CO514528 s13384_MA
13	17.2	78.2	463	1	AU083559
14	17.2	78.2	479	1	AU089685
15	17.2	78.2	513	4	BJ094274
16	17.2	78.2	515	7	CF447937
17	17.2	78.2	519	4	BI796581
18	17.2	78.2	571	4	BM037907
19	17.2	78.2	595	9	CC952473
20	17.2	78.2	617	6	CD488495
21	17.2	78.2	619	9	CL956886
22	17.2	78.2	633	4	BJ808940
23	17.2	78.2	634	7	CR286398
24	17.2	78.2	655	6	CD487922

25	17.2	78.2	726	5	BM071434
26	17.2	78.2	767	6	CB685128
27	17.2	78.2	812	6	CB644373
28	17.2	78.2	844	6	CB685127
29	17.2	78.2	852	9	CL670249
30	17.2	78.2	882	7	CF378583
31	17.2	78.2	972	6	CNS05PD9
32	17.2	78.2	1028	6	CA139194
33	17.2	78.2	1852	9	CL487297
34	17.2	78.2	2481	3	AK037625
35	16.8	76.4	105	1	AA094019
36	16.8	76.4	496	9	CE537167
37	16.8	76.4	523	8	AZ483488
38	16.8	76.4	526	8	AZ501799
39	16.8	76.4	628	6	CA380211
40	16.8	76.4	645	6	CB576172
41	16.8	76.4	654	7	CO079691
42	16.8	76.4	678	6	CA373611
43	16.8	76.4	681	1	AV732648
44	16.8	76.4	683	7	CV510488
45	16.8	76.4	705	2	AW916461

## ALIGNMENTS

RESULT 1  
BH859011  
LOCUS BH859011 521 bp DNA linear GSS 13-NOV-2002  
DEFINITION S5\_182b t7 Mouse Retroviral Tagged Cancer Gene Database Mus  
musculus genomic clone S5\_182b, genomic survey sequence.  
ACCESSION BH859011 GI:21709832  
VERSION BH859011.1  
KEYWORDS GSS.  
SOURCE Mus musculus (house mouse)  
ORGANISM Mus musculus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
REFERENCE 1 (bases 1 to 521)  
AUTHORS Suzuki, T., Shen, H., Akagi, K., Morse, H.C., Malley, J.D., Naiman, D.Q.,  
Jenkins, N.A. and Copeland, N.G.  
TITLE New genes involved in cancer identified by retroviral tagging  
JOURNAL Nat. Genet. 32 (1), 166-174 (2002)  
MEDLINE 22194816  
PUBMED 12185365  
COMMENT Contact: Copeland NG  
Mouse Cancer Genetics Program  
National Cancer Institute  
Bldg. 539, Rm. 229, Frederick, MD 21702-1201, USA  
Tel: 301 846 1260  
Fax: 301 846 6566  
Email: copeland@ncifcrf.gov  
Classes: PCR with specific primers.  
Location/Qualifiers  
1. .521  
/organism="Mus musculus"  
/mol\_type="genomic DNA"  
/db\_xref="taxon:10090"  
/clone="S5\_182b"  
/sex="female"  
/tissue\_type="leukemia"  
/note="Inverse PCR method  
(http://genome2.ncifcrf.gov/RTCGD)"

## FEATURES

source  
1. .521  
/organism="Mus musculus"  
/mol\_type="genomic DNA"  
/db\_xref="taxon:10090"  
/clone="S5\_182b"  
/sex="female"  
/tissue\_type="leukemia"  
/note="Inverse PCR method  
(http://genome2.ncifcrf.gov/RTCGD)"

## ORIGIN

Query Match 85.5%; Score 18.8; DB 8; Length 521;  
Best Local Similarity 90.9%; Pred. No. 91;  
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
QY 1 TGACTGTGAACGTTTCGAGATGA 22  
|||||  
DB 116 TGACTGTGAACATCGGAGATGA 137  
|||||

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RESULT 2
LOCUS
DEFINITION
BM042508          571 bp    mRNA    linear    EST 07-NOV-2001
603615795Tt NTH_MGC_112 Homo sapiens cDNA clone IMAGE:5420734 3',
mRNA sequence.
ACCESSION
BM042508
VERSION
BM042508.1 GI:16771788
KEYWORDS
EST.
SOURCE
Homo sapiens (human)
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1. (bases 1 to 571)
/organism="Homo sapiens"
AUTHORS
NIH-MGC http://mgc.nci.nih.gov/.
TITLE
National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL
Unpublished (1999)
COMMENT
Contact: Robert Strausberg, Ph.D.
Email: cgabbs@mail.nih.gov
Tissue Procurement: DCTD/DTP
cDNA Library Preparation: Ling Hong/Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone Distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLC1875 row: m column: 23
High quality sequence start: 44
High quality sequence stop: 411.
Location/Qualifiers
1. .571
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:5420734"
/tissue_type="melanotic melanoma, cell line"
/lab_host="DH10B (phage-resistant)"
/clone_lib="NIH MGC_112"
/note="Organ: Skin; Vector: pOTB7; Site 1: XhoI; Site 2:
EcoRI; cDNA made by oligo-dT priming. Directionally cloned
into EcoRI/XhoI sites using the following 5' adaptor:
GGCAGCAG(G). Library constructed by Ling Hong in the
laboratory of Gerald M. Rubin (University of California,
Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and
Superscript II RT (Life Technologies). Note: this is a
NIH_MGC Library."
ORIGIN
Query Match 83.6%; Score 18.4; DB 4; Length 571;
Best Local Similarity 95.0%; Pred. No. 1.5e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTTCGAGAT 20
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Db 504 TGACTGTGAACGTTTCAGAT 523
|||||

RESULT 3
CE751403/c
LOCUS
DEFINITION
tigr-gss-dog-17000369615400 Dog Library Canis familiaris genomic,
genomic survey sequence.
ACCESSION
CE751403
VERSION
CE751403.1 GI:37092020
KEYWORDS
GSS.
SOURCE
Canis familiaris (dog)
ORGANISM
Canis familiaris
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Carnivora; Pissipedia; Canidae; Canis.
REFERENCE
1. (bases 1 to 492)
/organism="Canis familiaris"
/db_xref="taxon:9606"
/clone="ev02g09"
/sex="Male"
/lab_host="DH10B"
/lab_lib="PAX3 CASTING Library 'ev'"
Rusch,D.B., Deicher,A.L., Pop,M., Wang,W., Fraser,C.M. and
Kirkness,E.F., Batna,V., Halpern,A.L., Levy,S., Remington,K.,
Venter,J.C.
The dog genome: survey sequencing and comparative analysis
Science 301 (5641), 1898-1903 (2003)
22875432
MEDLINE
PUBMED
14512627
Contact: Kirkness EF
The Institute for Genomic Research
Department of Eukaryotic Genomics, TIGR, 9712 Medical Center Drive,
Rockville, MD 20850, USA
Tel: 301-838-0200
Fax: 301-838-0208
Email: ekirknes@tigr.org
Class: shotgun.
Location/Qualifiers
1. .492
/organism="Canis familiaris"
/mol_type="genomic DNA"
/strain="Standard Poodle"
/db_xref="taxon:9615"
/clone_lib="Dog Library"
/note="Site 1: BstXI; Libraries were prepared from
peripheral blood"
ORIGIN
Query Match 80.9%; Score 17.8; DB 9; Length 492;
Best Local Similarity 90.5%; Pred. No. 3e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTTCGAGATG 21
|||||
Db 36 TGACTGTGAACGTCGCGGATG 16
|||||

RESULT 4
AZ755668/c
LOCUS
DEFINITION
ev02g09.x1 PAX3 CASTING Library 'ev' Homo sapiens genomic clone
ev02g09 random, genomic survey sequence.
ACCESSION
AZ755668
VERSION
AZ755668.1 GI:13175090
KEYWORDS
GSS.
SOURCE
Homo sapiens (human)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1. (bases 1 to 561)
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="ev02g09"
/lab_host="NIH MGC_112"
/note="Organ: Skin; Vector: pOTB7; Site 1: XhoI; Site 2:
EcoRI; cDNA made by oligo-dT priming. Directionally cloned
into EcoRI/XhoI sites using the following 5' adaptor:
GGCAGCAG(G). Library constructed by Ling Hong in the
laboratory of Gerald M. Rubin (University of California,
Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and
Superscript II RT (Life Technologies). Note: this is a
NIH_MGC Library."
AUTHORS
Friedman,T.B.
Identification of Target Genes Regulated by PAX3 and PAX3--FKHR in
Embryogenesis and Alveolar Rhabdomyosarcoma
Genomics 79 (3), 278-284 (2002)
21853298
MEDLINE
PUBMED
11863357
Contact: Friedman TB
Laboratory of Molecular Genetics
National Institute on Deafness and Other Communication Disorders,
National Institutes of Health
5 Research Court, Room 2A-15, Rockville, MD 20850, USA
Tel: 301 402 7580
Fax: 301 496 7882
Email: friedman@nidcd.nih.gov
Plate: 02 row: g column: 09
Seq primer: -21M13 forward primer (ABI)
Class: random plasmid subclone.
Location/Qualifiers
1. .561
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/clone="ev02g09"
/sex="Male"
/lab_host="DH10B"
/lab_lib="PAX3 CASTING Library 'ev'"
FEATURES
source

```



/note="Vector: pGEM-T Easy; Human genomic DNA was partially digested with Sau3AI, ligated to ds linkers, and enriched for binding to human PAX3DQ+ protein using a whole genome PCR-based strategy. DNA fragments containing putative PAX3DQ+ binding sites were amplified by PCR and cloned into pGEM-T Easy (Promega). The ligation products were transformed into DH10B electrocompetent cells (Life Technologies)."

## ORIGIN

Query Match 80.9%; Score 17.8; DB 8; Length 561;  
Best Local Similarity 90.5%; Pred. No. 3.1e+02;  
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTTCGAGATG 21  
|||||  
Db 461 TGACTGTGAACGTTTCGAGATG 441

## RESULT 5

BZ558601 867 bp DNA linear GSS 17-DEC-2002  
LOCUS pa98401\_292.s1 pac52-164 Pseudomonas aeruginosa genomic clone  
DEFINITION pa98401\_292, genomic survey sequence.

ACCESSION BZ558601.1 GI:27173329

## VERSION

KEYWORDS GSS.

## SOURCE

ORGANISM Pseudomonas aeruginosa

Bacteria; Proteobacteria; Gammaproteobacteria; Pseudomonadales;

Pseudomonadaceae; Pseudomonas.

1 (bases 1 to 867)

Spencer, D.H., Raymond, C.K., Smith, E.E., Sims, E.E., Hastings, M.,

Burns, J.L., Kaul, R., and Olsen, M.V.

Whole-Genome-Sequence variation among multiple isolates of

Pseudomonas aeruginosa library

J. Bacteriol. (2002) In press

Contact: Chris K. Raymond

Genome Center

University of Washington

Box 352145, Seattle, WA 98105-2145, USA

Tel: 2062216954

Fax: 2066857244

Email: craymond@u.washington.edu

Class: shotgun.

## FEATURES

## source

1. .867

/organism="Pseudomonas aeruginosa"

/mol\_type="genomic DNA"

/strain="2-164"

/db\_xref="taxon:287"

/clone="pa98401\_292"

/clone\_lib="pac52-164"

/note="Clinical isolate 2-164 Whole genomic shotgun

library."

## ORIGIN

Query Match 80.9%; Score 17.8; DB 8; Length 867;  
Best Local Similarity 90.5%; Pred. No. 3.3e+02;  
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GACTGTGAACGTTTCGAGATGA 22  
|||||  
Db 724 GACTGTGAACGTTTCGATATGA 744

## RESULT 6

BF971856 961 bp mRNA linear EST 22-JAN-2001  
LOCUS 602240444F1 NIH\_MGC\_46 Homo sapiens cDNA clone IMAGE:4328890 5',  
DEFINITION mRNA sequence.

## ACCESSION

VERSION BF971856.1 GI:12339071

## KEYWORDS

## SOURCE

## ORGANISM

## REFERENCE

## AUTHORS

## TITLE

## JOURNAL

## COMMENT

EST. Homo sapiens (human)

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 961)

NIH-MGC <http://mgs.nci.nih.gov/>

National Institutes of Health, Mammalian Gene Collection (MGC)

Unpublished (1999)

Contact: Robert Strausberg, Ph.D.

Email: [cgabbs@mail.nih.gov](mailto:cgabbs@mail.nih.gov)

Tissue Procurement: ATCC

CDNA Library Preparation: Ling Hong/Rubin Laboratory

DNA Sequencing by: Incyte Genomics, Inc.

Clone distribution: MGC clone distribution information can be

found through the I.M.A.G.E. Consortium/LNL at:

<http://image.lnl.gov>

Plate: LLCM189 row: h column: 11

High quality sequence stop: 555.

Location/Qualifiers

1. .961

/organism="Homo sapiens"

/mol\_type="mRNA"

/db\_xref="taxon:9606"

/clone="IMAGE:4328890"

/tissue\_type="leiomyosarcoma cell line"

/lab\_host="DH10B (phage-resistant)"

/clone\_lib="NIH MGC 46"

/note="Organ: uterus; Vector: pOTB7; Site: 1: XhoI; Site 2:

EcoRI; cDNA made by oligo-dT priming. Directionally cloned

into EcoRI/XhoI sites using the following 5' adaptor:

GCACACAG(G). Size-selected >500bp for average insert size

1.8kb. Library constructed by Ling Hong in the laboratory

of Gerald M. Rubin (University of California, Berkeley)

using ZAP-cDNA synthesis kit (Stratagene) and Superscript

II RT (Life Technologies). Note: this is a NIH\_MGC

Library."

## ORIGIN

Query Match 80.9%; Score 17.8; DB 4; Length 961;  
Best Local Similarity 90.5%; Pred. No. 3.4e+02;  
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GACTGTGAACGTTTCGAGATGA 22  
|||||  
Db 650 GACTGTGAACGTTTCGAGATGA 670

## RESULT 7

## LOCUS

## DEFINITION

## ACCESSION

## VERSION

## KEYWORDS

## SOURCE

## ORGANISM

## REFERENCE

## AUTHORS

## TITLE

## JOURNAL

## COMMENT

AZ060178 489 bp DNA linear GSS 30-MAR-2000  
RPCI-23-405E23.TJ RPCI-23 Mus musculus genomic clone  
RPCI-23-405E23, genomic survey sequence.

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

AZ060178 489 bp DNA linear GSS 30-MAR-2000

```

Fax: 301 838 0208
Email: szhao@tigr.org
Clones are derived from the mouse BAC library RPCI-23. For BAC
library availability, please contact Pieter de Jong
(piet@dejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buffalo.edu/orderingframe.htm)
or from Resea ch Genetics (info@resgen.com). BAC end page:
http://www.tigr.org/tdb/bac_ends/mouse/bac_end_intro.html
Plate: 405 row: E column: 23
Seq primer: SP6
Class: BAC ends.

FEATURES
    source
        1..489
            Location/Qualifiers
                /organism="Mus musculus"
                /mol_type="genomic DNA"
                /strain="C57BL/6J"
                /db_xref="taxon:10090"
                /clone="RPCI-23-405E23"
                /sex="Female"
                /lab_host="DH10B"
                /clone_lib="RPCI-23"
                /note="Organ: Kidney/Brain; Vector: pBACe3.6; Site_1:
                ECoRI; Site_2: ECoRI; Female C57BL/6J mouse kidney and/or
                brain genomic DNA was isolated and partially digested
                with a combination of ECoRI and EcoRI Methylase. Size
                selected DNA was cloned into the pBACe3.6 vector at the
                EcoRI sites. The ligation products were transformed into
                DH10B electrocompetent cells (BRL Life Technologies)."
    ORIGIN
        Query Match          79.1%; Score 17.4; DB 8; Length 489;
        Best Local Similarity 94.7%; Pred. No. 4.9e+02;
        Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

    QY 1 TGACTGTGAACGTTTCGAGA 19
        |||||||
    Db 170 TGACTGTGAACATTCGAGA 152
        |||||||

RESULT 8
AZ886419/c
LOCUS
DEFINITION
    AZ886419 530 bp DNA linear GSS 05-MAR-2001
    genomic survey sequence.
ACCESSION
AZ886419.1 GI:13205364
VERSION
GSS.
SOURCE
Mus musculus (house mouse)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
1 (bases 1 to 530)
Zhao,S., Nierman,W., Feldblyum,T., Malek,J., Shatsman,S.,
Akinret,B., Levine,M., McGann,S., Tsegaye,G., Geer,K., Krol,M., de
Jong,P. and Fraser,C.M.
Mouse BAC End Sequences from Library RPCI-23
Unpublished (1999)
TITLE
Mouse BAC End Sequences from Library RPCI-23
JOURNAL
COMMENT
Other GSSs: RPCI-23-18216.TV
Contact: Shaying Zhao
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208
Email: szhao@tigr.org
Clones are derived from the mouse BAC library RPCI-23. For BAC
library availability, please contact Pieter de Jong
(pdejong@mail.cho.org). Clones may be purchased from BACPAC
Resources (http://www.choi.org/bacpac/orderingframe.htm). BAC end
page: http://ww.tigr.org/tdb/bac_ends/mouse/bac_end_intro.html
Plate: 182 row: I column: 6
Seq primer: SP6
Class: BAC ends.

FEATURES
    source
        1..374
            Location/Qualifiers
                /organism="Homo sapiens"
                /mol_type="genomic DNA"
                /db_xref="taxon:9606"
                /clone="Plate=2056 Col=5 Row=J"
                /sex="male"
                /clone_lib="CIT Approved Human Genomic Sperm Library D"
                /note="Organ: sperm; Vector: pBeloBAC11; BAC Clones in
                E-Coli DH10B"
    ORIGIN
        Query Match          79.1%; Score 17.4; DB 8; Length 530;
        Best Local Similarity 94.7%; Pred. No. 4.9e+02;
        Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

    QY 1 TGACTGTGAACGTTTCGAGA 19
        |||||||
    Db 189 TGACTGTGAACATTCGAGA 171
        |||||||

RESULT 9
AQ245026
LOCUS
DEFINITION
    HS 2056 B1 E03 MR CIT Approved Human Genomic Sperm Library D Homo
    sapiens genomic clone Plate=2056 Col=5 Row=J, genomic survey
    sequence.
ACCESSION
AQ245026
VERSION
AQ245026.1 GI:3691600
KEYWORDS
GSS.
SOURCE
Homo sapiens (human)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 374)
Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
Hood,L.
Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
JOURNAL
MEDLINE
PUBMED
10449764
COMMENT
Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Sequence Tagged Connector
Plate: 2056 row: J column: 5
Class: BAC ends
High quality sequence stop: 374.

FEATURES
    source
        1..374
            Location/Qualifiers
                /organism="Homo sapiens"
                /mol_type="genomic DNA"
                /db_xref="taxon:9606"
                /clone="Plate=2056 Col=5 Row=J"
                /sex="male"
                /clone_lib="CIT Approved Human Genomic Sperm Library D"
                /note="Organ: sperm; Vector: pBeloBAC11; BAC Clones in
                E-Coli DH10B"
    ORIGIN

```

Query Match 78.2%; Score 17.2; DB 8; Length 374;  
 Best Local Similarity 86.4%; Pred. No. 5.9e+02;  
 Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTTCGAGATGA 22  
 |||||  
 Db 207 TGACTGTGAACGATTGAGATCA 228  
 |||||

RESULT 10  
 AZ536502  
 LOCUS  
 DEFINITION 110300\_96 Planococcus lilacinus DNA Planococcus lilacinus genomic,  
 genomic survey sequence.  
 ACCESSION AZ536502  
 VERSION AZ536502.1 GI:11093449  
 KEYWORDS GSS.  
 SOURCE Planococcus lilacinus (lilac mealybug)  
 ORGANISM Planococcus lilacinus  
 Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;  
 Neoptera; Paraneoptera; Hemiptera; Sternorrhyncha; Aphidiformes;  
 Coccoidea; Pseudococcidae; Planococcus.  
 REFERENCE 1 (bases 1 to 408)  
 AUTHORS Mohan, K.N. and Chandra, H.S.  
 TITLE Mealybug shotgun sequencing  
 JOURNAL Unpublished (2000)  
 COMMENT Contact: Mohan KN  
 Indian Institute of Science  
 Sir C.V. Raman Avenue, Bangalore, Karnataka 560012, India  
 Email: mohan@cbl.iisc.ernet.in  
 Class: shotgun.

FEATURES  
 source  
 1. .408  
 Location/Qualifiers  
 /organism="Planococcus lilacinus"  
 /mol\_type="genomic DNA"  
 /db\_xref="taxon:40930"  
 /clone\_lib="Planococcus lilacinus DNA"

ORIGIN  
 Query Match 78.2%; Score 17.2; DB 8; Length 408;  
 Best Local Similarity 86.4%; Pred. No. 6e+02;  
 Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTTCGAGATGA 22  
 |||||  
 Db 251 TGACTGTGAACGACGATATGA 272  
 |||||

RESULT 11  
 BE723539  
 LOCUS  
 DEFINITION 193384 MARC 4BOV Bos taurus cDNA 5', mRNA sequence.  
 ACCESSION BE723539  
 VERSION BE723539.1 GI:10124826  
 KEYWORDS EST.  
 SOURCE Bos taurus (cow)  
 ORGANISM Bos taurus  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;  
 Bovinae; Bos.  
 REFERENCE 1 (bases 1 to 424)  
 AUTHORS Smith, T.P.L., Grosse, W.M., Freking, B.A., Roberts, A.J., Stone, R.T.,  
 Casas, E., Wray, J.E., White, J., Cho, J., Fahrenkrug, S.C.,  
 Bennett, G.L., Heaton, M.P., Laegreid, W.W., Rohrer, G.A.,  
 Chitko-McKown, C.G., Perte, G., Holt, I., Karanycheva, S., Liang, F.,  
 Quackenbush, J. and Keele, J.W.  
 TITLE Sequence evaluation of four pooled-tissue normalized bovine cDNA  
 libraries and construction of a gene index for cattle  
 JOURNAL Genome Res. 11 (4), 626-630 (2001)  
 MEDLINE 21180013  
 PubMed 11282978  
 COMMENT Contact: Smith TPL

USDA, ARS, US Meat Animal Research Center  
 PO Box 166, Clay Center, NE 68933-0166, USA  
 Tel: 402 762 4366  
 Fax: 402 762 4390  
 Email: smith@email.marc.usda.gov  
 Single pass sequencing. Bases called and alt trimmed with phred  
 v0.980904.e. Vector identified by cross\_match with the -minscore 18  
 and -minmatch 12 options.  
 PCR Primers  
 FORWARD: AGGAAACAGCTATGACCAT  
 BACKWARD: GTTTCGCCAGTCACGACG  
 Plate: 92 row: B column: 14  
 Seq primer: ATTAGTGACACTATAG.  
 Location/Qualifiers  
 1. .424  
 /organism="Bos taurus"  
 /mol\_type="mRNA"  
 /db\_xref="taxon:9913"  
 /tissue\_type="pooled"  
 /lab\_host="DH10B"  
 /clone\_lib="MARC 4BOV"  
 /note="Vector: pCMV SPORT6; Site 1: NotI; Site 2: SalI;  
 Library made from pooled tissue from day 20 and day 40  
 embryos."

ORIGIN  
 Query Match 78.2%; Score 17.2; DB 2; Length 424;  
 Best Local Similarity 86.4%; Pred. No. 6e+02;  
 Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTTCGAGATGA 22  
 |||||  
 Db 268 TGAAGTCTGAACGTTAGAGATGA 289  
 |||||

RESULT 12  
 COS14528/c  
 LOCUS  
 DEFINITION 813DSG43G0800066\_327716 Glandular trichomes Medicago sativa cDNA,  
 mRNA sequence.  
 ACCESSION COS14528  
 VERSION COS14528.1 GI:50319402  
 KEYWORDS EST.  
 SOURCE Medicago sativa  
 ORGANISM Medicago sativa  
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;  
 rosids; eurosids I; Fabales; Fabaceae; Papilionoideae; Trifolieae;  
 Medicago.  
 REFERENCE 1 (bases 1 to 427)  
 AUTHORS Aziz, N., May, G.D., Paiva, N.L. and Dixon, R.A.  
 TITLE Alfalfa trichome Expressed Sequence Tags from the Samuel Roberts  
 Noble Foundation - Center for Medicago Genomics Research  
 JOURNAL Unpublished (2004)  
 COMMENT Contact: May GD  
 Plant Biology Division  
 The Samuel Roberts Noble Foundation  
 2510 Sam Noble Parkway, Ardmore, OK 73402, USA  
 Tel: 580 224 6650  
 Fax: 580 224 6692  
 Email: gdmay@noble.org.

FEATURES  
 source  
 1. .427  
 Location/Qualifiers  
 /organism="Medicago sativa"  
 /mol\_type="mRNA"  
 /db\_xref="taxon:3879"  
 /tissue\_type="Glandular trichomes isolated from stem"  
 /dev\_stage="Trichomes were removed from internodes of 8-12  
 inch tall stems"  
 /clone\_lib="Glandular trichomes"  
 /note="Vector: pDNR-LJB; Glandular-haired alfalfa plants  
 were established in a Conviron growth chamber (16-h days,  
 full lights, 24oc set point) in 40 one gallon pots. Plants

were grown in Metromix 350 and fertilized with MiracleGro as needed. They were cut back closely to encourage the emergence of vigorous shoots. Trichomes were isolated from stems, approximately 8-12 inches long, clipped from plants 2-3 inches above the crown. With minimal handling of the stem, the apical bud, leaves and nodes were discarded, and the trichomes isolated from the internode segments."

## ORIGIN

Query Match 78.2%; Score 17.2; DB 7; Length 427;  
Best Local Similarity 86.4%; Pred. No. 6e+02;  
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTTCGAGATGA 22

Db 198 TGATTGTGAACCTTCAGATGA 177

## RESULT 13

AU083559

LOCUS AU083559 463 bp mRNA linear EST 02-APR-2002  
DEFINITION AU083559 Rice green shoot Oryza sativa (japonica cultivar-group)  
CDNA clone S14862, mRNA sequence.

ACCESSION AU083559.1 GI:7274015

VERSION

KEYWORDS

SOURCE

ORGANISM

Oryza sativa (japonica cultivar-group)  
Oryza sativa (japonica cultivar-group)  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;  
Ehrhartoideae; Oryzeae; Oryza.

1 (bases 1 to 463)

Sasaki, T. and Yamamoto, K.

Rice cDNA from green shoot (2000)

Unpublished (2000)

Contact: Takuji Sasaki

National Institute of Agrobiological Resources

Rice Genome Research Program, Kannondai 2-1-2, Tsukuba, Ibaraki

305-8602, Japan

Tel: 81-298-38-7441

Fax: 81-298-38-7468

Email: tsasaki@nri.affrc.go.jp, URL: http://rgp.dna.affrc.go.jp/

PROJECT = 'RGP'

## FEATURES

source

1..463

/organism="Oryza sativa (japonica cultivar-group)"

/mol\_type="mRNA"

/cultivar="Nipponbare"

/db\_xref="taxon:39947"

/clone="S14862"

/clone\_lib="Rice green shoot"

/notes="Green shoot (8 days old)"

## ORIGIN

Query Match 78.2%; Score 17.2; DB 1; Length 463;  
Best Local Similarity 86.4%; Pred. No. 6.1e+02;  
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTTCGAGATGA 22

Db 151 TGAGTGTGAATGTTAGATGA 172

## RESULT 14

AU089685

LOCUS

DEFINITION AU089685 Rice callus Oryza sativa (japonica cultivar-group) cDNA  
clone C40060, mRNA sequence.

ACCESSION AU089685

VERSION

KEYWORDS

SOURCE

ORGANISM

Oryza sativa (japonica cultivar-group)

Oryza sativa (japonica cultivar-group)

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;  
Ehrhartoideae; Oryzeae; Oryza.

1 (bases 1 to 479)

Sasaki, T. and Yamamoto, K.

Rice cDNA from callus (2000)

Unpublished (2000)

Contact: Takuji Sasaki

National Institute of Agrobiological Resources

Rice Genome Research Program, Kannondai 2-1-2, Tsukuba, Ibaraki

305-8602, Japan

Tel: 81-298-38-7441

Fax: 81-298-38-7468

Email: tsasaki@nri.affrc.go.jp, URL: http://rgp.dna.affrc.go.jp/

PROJECT = 'RGP'

## FEATURES

source

1..479

/organism="Oryza sativa (japonica cultivar-group)"

/mol\_type="mRNA"

/cultivar="Nipponbare"

/db\_xref="taxon:39947"

/clone="C40060"

/clone\_lib="Rice callus"

/note="Vector: pBluescript II SK+; Site 1: SalI; Site 2:

NotI; cDNA prepared from rice callus mRNAs by using

oligo(dT) as a primer and ligating to the SalI-NotI site

of pBluescript II SK+ phagemid."

## ORIGIN

Query Match 78.2%; Score 17.2; DB 1; Length 479;  
Best Local Similarity 86.4%; Pred. No. 6.2e+02;  
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTTCGAGATGA 22

Db 299 TGAGTGTGAATGTTAGATGA 320

## RESULT 15

BU094274

LOCUS

DEFINITION BU094274 NTBB Mochii normalized Xenopus early gastrula library  
Xenopus laevis cDNA clone XL144p18 5', mRNA sequence.

ACCESSION BU094274

VERSION

KEYWORDS

SOURCE

ORGANISM

Xenopus laevis (African clawed frog)

Xenopus laevis

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Amphibia; Batrachia; Anura; Mesobatrachia; Pipidae;

Xenopodinae; Xenopus; Xenopus.

1 (bases 1 to 513)

Kitayama, A., Terasaka, C., Mochii, M., Ueno, N., Shin-i, T. and

Kohara, Y.

Expressed genes in X. laevis embryo

Unpublished (2001)

Contact: Tadasu Shin-i

Center For Genetic Resource Information

National Institute of Genetics

1111 Yata, Mishima, Shizuoka 411-8540, Japan

Tel: 81-559-81-6856

Fax: 81-559-81-6855

Email: tshini@genes.nig.ac.jp

The information of this clone is available through the following

URL.

http://xenopus.nibb.ac.jp.

## FEATURES

source

1..513

/organism="Xenopus laevis"

/mol\_type="mRNA"

/db\_xref="taxon:8355"

/clone="XL144p18"

/tissue\_type="whole embryo"

U.S. National Library of Medicine

## ORIGIN

Best Local Similarity 86.4%; Pred: No. 6.2e+02;  
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 TGACTGTGAACGTTTCGAGATGA 22

Db  
198 TGCCTGAGAACGTTTCGCGATGA 219

Job time : 1998 secs

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